

GenCore version 5.1.3  
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OM Protein - Protein search, using sw model

Run on: February 20, 2003, 10:46:10 ; Search time 12 seconds

(without alignments)  
48.067 Million cell updates/sec

Title: US-09-912-414-9

Perfect score: 31

Sequence: 1 WXXWVF 6

Scoring table: BLOSUM62PX

Gapop: 10.0, Gapext: 0.5

Searched: 283224 seqs, 9634422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-Processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database:

PIR\_73: \*  
1: p1r1: \*  
2: p1r2: \*  
3: p1r3: \*  
4: p1r4: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	31	100.0	21	2	PIR1731
2	31	100.0	49	2	I48681
3	31	100.0	59	2	E82844
4	31	100.0	61	2	H91196
5	31	100.0	65	2	S23164
6	31	100.0	71	2	F84168
7	31	100.0	73	2	A82788
8	31	100.0	75	2	T14724
9	31	100.0	78	2	S84446
10	31	100.0	79	2	A64750
11	31	100.0	83	2	A97568
12	31	100.0	86	2	T42185
13	31	100.0	87	2	H86753
14	31	100.0	92	2	S75609
15	31	100.0	92	2	TG1637
16	31	100.0	93	2	G81098
17	31	100.0	97	2	E84147
18	31	100.0	99	2	G84022
19	31	100.0	101	2	S75818
20	31	100.0	103	2	F82266
21	31	100.0	105	2	J82384
22	31	100.0	106	2	T64171
23	31	100.0	106	2	G82715
24	31	100.0	106	2	G72912
25	31	100.0	108	2	A91164
26	31	100.0	108	2	S28241
27	31	100.0	108	2	A93505
28	31	100.0	109	2	F83057
29	31	100.0	109	2	A72029

30	31	100.0	110	2	AD0127	conserved hypotet
31	31	100.0	110	2	AG1523	hypothetical prote
32	31	100.0	116	2	P89666	Ig heavy chain pre
33	31	100.0	116	2	G82398	conserved hypotet
34	31	100.0	117	2	A95381	hypothetical prote
35	31	100.0	118	2	S24527	Ig heavy chain V r
36	31	100.0	119	2	S24522	Ig heavy chain V r
37	31	100.0	119	2	S24492	Ig heavy chain V r
38	31	100.0	119	2	S24513	Ig heavy chain V r
39	31	100.0	119	2	S24493	Ig heavy chain V r
40	31	100.0	119	2	S24491	Ig heavy chain V r
41	31	100.0	119	2	S24525	Ig heavy chain V r
42	31	100.0	119	2	S24523	Ig heavy chain V r
43	31	100.0	119	2	S24518	Ig heavy chain V r
44	31	100.0	119	2	S24511	Ig heavy chain V r
45	31	100.0	119	2	S24494	Ig heavy chain V r

#### ALIGNMENTS

##### RESULT 1

PIR1731

Ig heavy chain V region (clone GCC-11) - mouse (fragment)

C/Species: Mus musculus (house mouse)

C/Date: 24-Feb-1994 #sequence\_revision 24 Feb 1994 #text\_change 17-Mar-1999

C/Accession: PIR1731

P.Melhyzer Williams, M.G., McLean, M.J., Lalor, F.A., Nossal, G.J.V.

U. Exp. Med. 178, 295-307, 1993

A>Title: Antigen-driven B cell differentiation in vivo.

A/Reference number: PIR1675; MIMD-93301607; PMID-831585

A/Accession: PIR1731

A/Molecule type: mRNA

A/Residues: 121 <MGH>

A/Experimental source: B cell

A/Note: the authors translated the codon ACA for residue 13 as Ala

A/Superfamily: immunoglobulin V region, immunoglobulin heavy

C/Keywords: heterotetramer, immunoglobulin

##### Query Match

100.0%; Score 31; DB 2; Length 21;

Best Local Similarity 50.0%; Pred No 12402;

Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWVF 6  
DB 16 WTSWST 21

##### RESULT 2

I48681

185 kDa glycoprophosphoprotein - mouse (fragment)

C/Species: Mus musculus (house mouse)

C/Date: 22-Jul-1996 #sequence\_revision 22-Jul-1996 #text\_change 07-Jun-2002

C/Accession: I48681

P.White, M.B.; Hung, M.C.

Oncogene 7, 677-683, 1992

A>Title: Cloning and characterization of the mouse gene for

A/Reference number: I48681, MIMD-9228496; PMID-1348955

A/Accession: I48681

A/Status: preliminary; translated from GP/EMBL/DBD

A/Molecule type: DNA

A/Residues: 1-49 <RES>

A/Cross-references: EMBL.X66236, MIMD-9325186, FIDR.CAA46966.1, FID 3705187

C/Genes: neu

C/Keywords: ATP; phosphoprotein

Query Match 100.0%; Score 31; DB 2; Length 49;

Best Local Similarity 50.0%; Pred No 27402;

Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWVF 6  
DB 16 WTSWST 21

D <sub>0</sub>	3	W	D	E	N	z	z

A>Title The volume of the National Centre Engineer Association Conference - 568  
A-Efficient member A00-77-000000000

A>Title The volume of the National Centre Engineer Association Conference - 568  
A-Efficient member A00-77-000000000

A:Accession: AH2788  
 A:Status: Preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-77 <REN>  
 A:Cross-references: GB:AF008844; FIDN:AA14776.1; FID:G17749165; UNFEB:DH00186  
 A:Experimental source: strain GSE (Dugont)  
 C:Genetics:  
 A:Gene: AtU1727  
 A:Map position: circular chromosome

Query Match 100.0%; Score 31; DB 2; Length 73;  
 Best Local Similarity 50.0%; Pred. No. 4.1e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6  
 Db 13 WNLWLF 18

RESULT 8  
 T14724  
 Hypothetical protein Y1031 - Yersinia pestis plasmid pMT1

C:Species: Yersinia pestis  
 C:Date: 20-Sep-1999 #sequence\_revision 20 Sep 1999 #ext\_change 24-Sep-1999  
 C:Accession: T14724; T15006  
 R:Hu, P.; Elliott, J.; McTernan, P.; Skowronski, F.; Garner, J.; Kobayashi, A.; Carraro,  
 submitted to the EMBL Data Library, March 1999  
 A:Description: Structural organization of virulence determinants in three Yersinia pestis  
 A:Reference number: T16168  
 A:Accession: T14724  
 A:Status: preliminary; translated from GB/EMBL/DBP  
 A:Molecule type: DNA  
 A:Residues: 1-76 <HUP>

A:Cross-references: EMBL:AF034947; FID:G2096246; FID:G2096163; FIDN:AA013241;  
 R:Blindley, L.E.; Plano, G.V.; Rutland, V.; Mayhew, G.F.; Blatner, F.P.  
 Infect. Immun. 66, 5731-5742, 1998  
 A:Title: Complete DNA sequence and detailed analysis of the Yersinia pestis KIMS plasmid  
 A:Reference number: 219264; MIMD:00043849; PMID:9826348  
 A:Accession: T15006  
 A:Status: preliminary; translated from GB/EMBL/DBP  
 A:Molecule type: DNA  
 A:Residues: 1-76 <HUP>

A:Cross-references: EMBL:AF074611; FID:G1493003; FID:G1493009; FIDN:AA097749;  
 C:Genetics:  
 A:Gene: Y1091  
 A:Gene: plasmid pMT1

Query Match 100.0%; Score 31; DB 2; Length 76;  
 Best Local Similarity 50.0%; Pred. No. 4e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6  
 Db 50 WNLWLF 55

RESULT 9  
 S08449

Hypothetical protein 3 Spiroplasma virus 1  
 C:Species: Spiroplasma virus 1, spvi  
 C:Date: 30-Jun-1991 #sequence\_revision 30-Jun-1991 #ext\_change 07-Dec-1999  
 C:Accession: S08449  
 R:Renaudin, J.; Aulio, P.; Vignault, J.C.; Bove, J.M.  
 Nucleic Acids Res. 19, 1293, 1990  
 A:Title: Complete nucleotide sequence of the genome of Spiroplasma citri virus Spvi-RNA2  
 A:Reference number: S08447; MIMD:00047699; PMID:2329423  
 A:Accession: S08449  
 A:Status: translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-78 <REN>  
 A:Cross-references: EMBL:X11441; FID:G11903; FIDN:AA16777.1; FID:G104400  
 C:Genetics:  
 A:Genetic code: SGC3

Query Match 100.0%; Score 31; DB 2; Length 78;  
 Best Local Similarity 50.0%; Pred. No. 4.1e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6  
 Db 42 WNLWLF 47

RESULT 10  
 A64750

Hypothetical protein h0249 - Escherichia coli (strain K-12)  
 C:Species: Escherichia coli  
 C:Date: 12-Sep-1997 #sequence\_revision 17-Sep-1997 #ext\_change 01-Mar-2002  
 C:Accession: A64750  
 R:Plattner, F.F.; Plunkett III, G.; Rhotch, C.A.; Perna, N.T.; Rutland, V.; Riley, M.; Col,  
 A.; Rose, D.J.; Mau, B.; Shao, Y.  
 Science 277, 1453-1462, 1997  
 A:Title: The complete genome sequence of Escherichia coli K-12.  
 A:Reference number: A64720; MIMD:0042617; PMID:9278501  
 A:Accession: A64750  
 A:Status: nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-79 <BLAT>

A:Cross-references: GB:AF064133; GB:U00006; FID:G1493003; FIDN:AA16777.1; FID:G104400  
 A:Experimental source: strain K-12, substrain W1655

Query Match 100.0%; Score 31; DB 2; Length 79;  
 Best Local Similarity 50.0%; Pred. No. 4.1e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6  
 Db 50 WXXXNF 55

RESULT 11  
 A97568

Hypothetical protein ACP\_3173 (inferred) Agrobacterium tumefaciens (strain C58, Cere  
 C:Species: Agrobacterium tumefaciens  
 C:Date: 30-Sep-2001 #sequence\_revision 30 Sep 2001 #ext\_change 11-Jun-2002  
 C:Accession: A97568  
 R:Goodner, B.; Hinkle, G.; Gelling, S.; Miller, N.; Blanchard, M.; Ouellet, B.; Gelfman,  
 A.; Liu, F.; Wellam, C.; Allinger, M.; DeGutty, D.; Scott, C.; Tafas, C.; Mikulicz, H.;  
 Science 224, 2233-2239, 2001  
 A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tum  
 A:Reference number: A97559; PMID:11743194  
 A:Accession: A97568  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-83 <KUR>

A:Cross-references: GB:AE007669; FIDN:AA074404; FID:G1493003; UNFEB:DH00186  
 C:Genetics:  
 A:Gene: ACP\_3173  
 A:Map position: circular chromosome

Query Match 100.0%; Score 31; DB 2; Length 83;  
 Best Local Similarity 50.0%; Pred. No. 4.2e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXNF 6  
 Db 23 WNLWLF 28

RESULT 12  
 T42185

conserved hypothetical protein L7085 - Escherichia coli plasmid p0157  
 C:Species: Escherichia coli  
 C:Date: 03-Dec-1994 #sequence\_revision 03-Dec-1994 #ext\_change 03-Dec-1999  
 C:Accession: T42185  
 R:Rutland, V.; Shao, Y.; Perna, N.T.; Plunkett, G.; Settla, H.; Blatner, F.P.





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OM protein. Protein search, using a modified

Run on: February 20, 2003, 10:41:59, Session: 12060667 Section:

37.329 Million coll updates/sec

Perfect score: 31

Sequence: 1 WXXWXF 6

Scoring table: BLOSUM62DX

Gapop 10.0 , Gapext 0.5

112997 seqs, 41476328 residues

Total number of hits on isoging: 11,002

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post Processing: Minimum Match 08

Post processing:	Minimum Match	08
	Maximum Match	10

Maximum	Match	100%
1st best	45%	

CONFIDENTIAL

DATE: 11/11/2011

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution

## SUMMARIES

Result	Query	Match	Length	Pos	Definition
1	31	100.0	65	1	HA22_ECTH
2	31	100.0	79	1	WCA_SPV1
3	31	100.0	79	1	YEEF_ECOLI
4	31	100.0	84	1	ACHA_OPDS
5	31	100.0	84	1	ACHA_FELCA
6	31	100.0	90	1	YB11_ECOLI
7	31	100.0	101	1	YGS0_MYCPN
8	31	100.0	106	1	NIGM_HAEMN
9	31	100.0	106	1	YGGC_HAEMN
10	31	100.0	104	1	NJ3M_BRYVN
11	31	100.0	116	1	HV05_CARAV
12	31	100.0	128	1	MERT_STAUV
13	31	100.0	129	1	YV3H_HUMAN
14	31	100.0	129	1	YIM2_YEAST
15	31	100.0	134	1	YAG1_MYCPN
16	31	100.0	139	1	YB13_MYCPN
17	31	100.0	144	1	HY26_MOUSE
18	31	100.0	145	1	P137_AEPEP
19	31	100.0	146	1	YGS6_YEAST
20	31	100.0	147	1	YGS6_METTA
21	31	100.0	147	1	H4P1_ECOLI
22	31	100.0	157	1	YEG3_MYCPN
23	31	100.0	164	1	LSPA_ECOLI
24	31	100.0	165	1	DSHR_VIBAL
25	31	100.0	165	1	LSPA_ENTAE
26	31	100.0	165	1	VPH1_PHTSH
27	31	100.0	166	1	Y1RK_SSV1
28	31	100.0	170	1	LSPA_PSEPL
29	31	100.0	171	1	Y42A_MYCPN
30	31	100.0	172	1	YB03_MYCPN
31	31	100.0	176	1	VPL_BPR1
32	31	100.0	177	1	TN1E_HUMAN
33	31	100.0	182	1	VG17_BPMH

34	31	100.0	196	1	VP8 YEAST	P46989	Saccharomyces
35	31	100.0	206	1	VP6 MYCTD	P15693	Myxobolus
36	31	100.0	204	1	VP6 ECT1	F57627	Escherichia
37	31	100.0	204	1	VP6 ECT1	E37515	Escherichia
38	31	100.0	210	1	SGH YEAST	G02446	Saccharomyces
39	31	100.0	213	1	IF4E XENLA	F48937	Xenopus lae
40	31	100.0	215	1	IF4E APICLA	G77210	Apicomplexa
41	31	100.0	217	1	IF4E HUMAN	F07733	Homo sapiens
42	31	100.0	217	1	IF4E MOUSE	F00471	Mus musculus
43	31	100.0	217	1	IF4E PARIT	D09339	Paritaxillus
44	31	100.0	219	1	PSF2 MPTA	E67366	Mycobacterium
45	31	100.0	220	1	RP28 BPSPI	P03048	Bacteriophage

## ALIGNMENTS

ID	NAME	STANDARD	PRT	AA
1	LMHA2_ECTHL			
2	LMHA2_ECTHL			
3	LMHA2_ECTHL			
4	LMHA2_ECTHL			
5	LMHA2_ECTHL			
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9	LMHA2_ECTHL			
10	LMHA2_ECTHL			
11	LMHA2_ECTHL			
12	LMHA2_ECTHL			
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14	LMHA2_ECTHL			
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97	LMHA2_ECTHL			
98	LMHA2_ECTHL			
99	LMHA2_ECTHL			
100	LMHA2_ECTHL			

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Query Match: 100.0%; Score: 41; PP: 1; Length: 65;
Best Local Similarity: 50.0%; Pred. No.: 1.3e+02;
Matches: 3; Conservative: 3; Mismatches: 0; Indels: 0; Gaps: 0;
```

RESULT 2  
VG9\_SPV1H

```

ID      V09 SPV16 STANDARD          PRT         78 AA
AC      P15909
DT      01-APR-1990 (Rel. 14, Created)
DT      01-FEB-1996 (Rel. 33, Last sequence update)
DT      01-FEB-1996 (Rel. 33, Last annotation update)
FE      Gene 3 Protein
GN      3
CC      EPICOLAEMA VIRUS GP1 PROD B.
CC      VirusE, DATA SOURCE, F00011146, FLUENTLIFE.
CX      NCBI_TaxId=10954;
LN
RP      SEQUENCE FROM N.A.
RX      MEDLINE=99266793; PubMed=2104433;
SA      Remington J., Ailio P., Vignatelli C., Rowland M. I.
PT      "Complete nucleotide sequence of the gene 3 protein of Epicolaema virus."
PI      Virus SpV-PAD B."
RL      Nucleic Acids Res. 18:1293-1299(1990).
CC
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CC      entities requires a license agreement. See http://www.ebi.ac.uk/databases/
CC      or send an email to license@ebi.ac.uk.
CC
CC      EMBL: X0114; CA01572.1;
CC      Sequence from AA, acc1.wg, refseq=EPICOLAEMA_VIRUS_3,
CC
CY      1 WXXWVF 6
CY      11111
DB      42 WHIMDF 47

```

[illegible]

SO SEQUENCE 84 AA; 9833 MW; 230672P8060796R CPO64;

Query Match: 100.0%; Score 31; FR 1; Length 84;  
Best Local Similarity 50.0%; Pred No 1 75-02;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXP 6  
[:::]  
Db 63 WKHWF 68

## RESULT 5

ACHA PELICA STANDARD; PRT; 84 AA.

ID ACHA PELICA

PS4250;

DT 01-OCT-1996 (rel 34, Created)

DT 01-OCT-1996 (rel 34, Last sequence update)

DT 15-JUN-2002 (rel 41, Last annotation update)

DE Acetylcholine receptor protein, alpha chain (Fragment)

GN CHRNA1.

OS Felis silvestris catus (Cat).

OC Eukaryota; Metazoa; Chordata; Carnivora; Fissipedata; Feliidae; Felis.

OC Mammalia; Eutheria; Carnivora; Fissipedata; Feliidae; Felis.

NCBI\_TaxID=9685;

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=Muscle;

RA MEDLINE=95345091; PubMed=7619817;

RA Baraban D, Ovadia M, Kochva E, Fuchs S;

RT The binding site of the nicotinic acetylcholine receptor in animal

RT Biochemistry 14:9172-9176(1995).

CC - FUNCTION: AFTER BINDING ACETYLCHOLINE, THE ACHR PERFORMS AN

CC EXTENSIVE CHANNEL IN CONFORMATION THAT AFFECTS ALL SUBUNITS AND

CC LEADS TO OPENING OF AN ION-CONDUCTING CHANNEL ACROSS THE PLASMA

CC MEMBRANE.

CC - SUPRAT. PENTAMER OF TWO ALPHA CHAINS, AND ONE EACH OF THE BETA,

CC DELTA, AND GAMMA (IN IMMATURE MUSCLE) OR Epsilon (IN MATURE

CC MUSCLE) CHAINS.

CC - SPECIFIC LIPID LOCATION: Integral membrane protein (By similarity).

CC - SIMILARITY: RELATES TO THE LIGAND-GATED IONIC CHANNEL FAMILY.

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## RESULT 6

YPOL ECOLI

ID YPOL ECOLI

PS4985;

DT 16-OCT-2001 (rel 40, Created)

DT 16-OCT-2001 (rel 40, Last sequence update)

DT 15-JUN-2002 (rel 41, Last annotation update)

DE Hypothetical protein ypol.

GN YPOL OP R241.1

OS Escherichia coli.

OC Bacteria; Proteobacteria; Gamma subdivision; Enterobacteriaceae;

OC Escherichia.

OC NCBI\_TaxID=562;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=K12 / MG1655;

RA MEDLINE=97425617; PubMed=9278503;

RA Blument F, Plank G, Li T, Plank G A, Perts N T, Pajand V,

RA Pilloy M, Collado-Vides J, Gilmart J B, Pote C K, Mayhew G F,

RA Greger J, Davis N W, Kitzberger H A, Gordon M A, Rose D J,

RA Mau B, Shao Y;

RT The complete genome sequence of Escherichia coli K-12.

RL Science 277:1453-1474(1997).

RN [2]

RP IDENTIFICATION, AND CONCEPTUAL TRANSLATION.

RA Budd K E;

RL Unpublished observations (Jan-1999).

CC - SIMILARITY: STRONG TO E COLI YPOL

CC - CAUTION: THIS IS A CONCEPTUAL TRANSLATION A FRAMESHIFT HAD TO BE

CC CORRECTED IN POSITION 60 AND TWO IN FRAME STOP CODONS HAD TO BE

CC SUPPRESSED IN POSITIONS 9 AND 11.

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## RESULT 7

YGO5 MYCPN

ID YGO5 MYCPN

PS5147;

DT 16-OCT-2001 (rel 40, Created)

DT 16-OCT-2001 (rel 40, Last sequence update)

DT 15-JUN-2002 (rel 41, Last annotation update)

DE Hypothetical lipid-protein MPN650 precursor (E09-01101).

GN MPN650 OP MP192

OS Mycoplasma pneumoniae.

OC Bacteria; Firmicutes; Mollicutes; Mycoplasmatraceae; Mycoplasma.

OC NCBI\_TaxID=2104;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=ATCC 29342 / M129,

RA MEDLINE=97105885; PubMed=8948633;

RA Himmelfrich R, Hilbert H, Plagens H, Pitkl E, La B C,

RA Hermann R;

RT Complete sequence analysis of the genome of the bacterium Mycoplasma

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[illegible]

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ID      YA91 MYCPN      STANDARD;      PRT;      138 AA.
AC      P75602;
BT      16-OCT-2001 (Rel. 49, Created)
DT      16-OCT-2001 (Rel. 40, Last sequence update)
DE      16-OCT-2001 (Rel. 40, Last annotation update)
DE      Hypothetical protein MPN091 (P02_01138).
GN      MPN091 OR MP064.
OS      Mycoplasma pneumoniae.
OC      Bacteria; Firmicutes; Mollicutes; Mycoplasmataceae; Mycoplasma.
OX      NCBI_TaxID=2104;
RN      (1)
RP      SEQUENCE FROM N.A.
RC      STRAIN ATCC 2942 / M129.
RX      MEDLINE:97105885; PubMed:8948633;
RA      Himmelreich R., Hilbert H., Plagens H., Pirk E., Li B.-C.,
RA      Herrmann R.;
RT      "Complete sequence analysis of the genome of the bacterium Mycoplasma
RT      pneumoniae."
RI      Nucleic Acids Res. 24:4420-4449(1996).
CC      -1- SIMILARITY: ALMOST IDENTICAL TO M.PNEUMONIAE MPN13 AND MPN463.
CC
CC      This SWISS-PROT entry is copyright. It is produced through a collaboration
CC      between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC      the European Bioinformatics Institute. There are no restrictions on its
CC      use by non-profit institutions as long as its content is in no way
CC      modified and this statement is not removed. Usage by and for commercial
CC      entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC      or send an email to license@sib.ch).
CC
DR      EMBL: AE000008; AAB95711.1; -
KW      Hypothetical protein; Complete proteome.
SQ      SEQUENCE 138 AA; 15763 MW; 452380D428F42709 CRC64;

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Query Match      100.0%; Score 31; DB 1; Length 138;
Best local Similarity 50.0%; Prod.No. 2.6e+02;
Matched 3; Overlapping 3; Mismatches 0; Gaps 0,

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QY      1 WXXXP 6
       11111
Db      36 MSGWSE 41

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Search completed: February 20, 2003, 10:48:02  
Job time : 7.66667 secs

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GenCode Version 5.1.3  
Copyright 1999-2003 Cambridge U.K.

OM protein - protein search, using sw model

Run on February 20, 2003, 10:45:10 : Search time 21.133 seconds  
(without alignments)  
52.044 Million cell reads/sec

Title: US-09-912-414-9

Perfect score: 31

Sequence: 1 MXXMXF 6

Scoring matrix: BLASTNCX

Gapop: 10.0, Gapext: 0.5

Searched: 671580 seqs, 206647115 residues

Total number of hits satisfying chosen parameters: 671580

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post processing: Minimum Match 98

Maximum Match 100%

Listing first 45 summaries

Database:

SPREMBL\_21:  
1: sp.archaea:  
2: sp.bacteria:  
3: sp.fungi:  
4: sp.human:  
5: sp.invertebrate:  
6: sp.mammal:  
7: sp.mbr:  
8: sp.organelle:  
9: sp.phage:  
10: sp.plant:  
11: sp.rodent:  
12: sp.virus:  
13: sp.vertebrate:  
14: sp.unclassified:  
15: sp.virus:  
16: sp.bacteriophage:  
17: sp.archaea:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution

## SUMMARIES

Result No	Score	Query	Match length	PF	ID	Description
1	31	100.0	28	6	Q62821	Q62821 bubalus bub
2	31	100.0	31	8	Q64812	Q64812 bubalus bub
3	31	100.0	36	15	Q60507	Q60507 bubalus bub
4	31	100.0	48	11	Q63168	Q63168 bubalus bub
5	31	100.0	49	11	Q68615	Q68615 mus musculu
6	31	100.0	50	4	Q6NRC0	Q6NRC0 homo sapien
7	31	100.0	50	16	Q6PH20	Q6PH20 xylella fas
8	31	100.0	61	16	Q6S619	Q6S619 escherichia
9	31	100.0	64	4	Q6WWM9	Q6WWM9 homo sapien
10	31	100.0	70	2	Q6XEX4	Q6XEX4 lactobacilli
11	31	100.0	71	17	Q6XST4	Q6XST4 halobacteri
12	31	100.0	74	2	Q6S168	Q6S168 mycobacteri
13	31	100.0	76	16	Q68793	Q68793 yersinia pe
14	31	100.0	77	2	Q6S4M3	Q6S4M3 escherichia
15	31	100.0	79	4	Q6MTF7	Q6MTF7 escherichia
16	31	100.0	79	16	Q6TMR3	Q6TMR3 escherichia

17	31	100.0	80	16	Q6TMR3	Q6TMR3 escherichia
18	31	100.0	83	16	Q6U8M9	Q6U8M9 agrobacteri
19	31	100.0	86	2	Q6S619	Q6S619 escherichia
20	31	100.0	86	2	Q6S619	Q6S619 escherichia
21	31	100.0	86	10	Q6W1M2	Q6W1M2 solanum cha
22	31	100.0	87	16	Q6S619	Q6S619 escherichia
23	31	100.0	89	10	Q6W1M9	Q6W1M9 solanum pin
24	31	100.0	89	10	Q6W1M5	Q6W1M5 solanum pin
25	31	100.0	92	10	Q6W1M0	Q6W1M0 solanum pin
26	31	100.0	92	12	Q6S619	Q6S619 escherichia
27	31	100.0	93	10	Q6W1M1	Q6W1M1 solanum pin
28	31	100.0	93	16	Q6S619	Q6S619 escherichia
29	31	100.0	94	4	Q6W6T4	Q6W6T4 homo sapien
30	31	100.0	95	4	Q6S619	Q6S619 escherichia
31	31	100.0	97	16	Q6S619	Q6S619 escherichia
32	31	100.0	98	16	Q6S619	Q6S619 escherichia
33	31	100.0	98	16	Q6S619	Q6S619 escherichia
34	31	100.0	101	5	Q6S619	Q6S619 escherichia
35	31	100.0	101	10	Q6X1M2	Q6X1M2 solanum cha
36	31	100.0	103	16	Q6S619	Q6S619 escherichia
37	31	100.0	103	16	Q6X1M2	Q6X1M2 solanum cha
38	31	100.0	105	11	Q6C8U2	Q6C8U2 mus musculu
39	31	100.0	106	16	Q6NRC0	Q6NRC0 homo sapien
40	31	100.0	106	16	Q6S619	Q6S619 escherichia
41	31	100.0	106	16	Q6X1M7	Q6X1M7 solanum cha
42	31	100.0	106	16	Q6S619	Q6S619 escherichia
43	31	100.0	107	2	Q6S619	Q6S619 escherichia
44	31	100.0	108	16	Q6S619	Q6S619 escherichia
45	31	100.0	108	16	Q6S619	Q6S619 escherichia

## ALIGNMENTS

RESULT 1  
ID: Q62821  
AC: Q62821  
DT: 01-AUG-1998 (TREMURel, 07, last sequence update)  
DT: 01-AUG-1998 (TREMURel, 07, last sequence update)  
DT: 01-DEC-2001 (TREMURel, 19, last annotation update)  
DE: Lysosome (Fragment)  
OS: Bubalus bubalis (Domestic water buffalo)  
OC: Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Ox; NCBI\_TaxID=89462;  
RN: [1]  
RP: SEQUENCE FROM N.A.  
RC: TISSUE=BLUDD;  
PA: Das P., Tiwari G., Gary L.C.;  
RT: "Characterisation of lysosome encoding gene promoter in Bubalus  
RT: bubalis";  
RT: Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.  
PL: EMBL, AJ225012; CNA12342.1;  
DR: NON TER  
FT: NON TER  
SQ: SEQUENCE 28 AA; 3160 MW; 886856AF7D08E9D CFC64;

Query Match: 100.0% Score: 31, 16.6, Length: 28;  
Post local similarity: 10.0% Pred. No. 2.0e+00;  
Matches: 3; Conservative: 3; Mismatches: 0; Indels: 0; Gaps: 0;

QY: 1 MXXMXF 6  
DB: 10 MSWMTF 15

RESULT 2  
ID: Q64812  
AC: Q64812  
DT: 01-DEC-2001 (TREMURel, 19, last sequence update)  
DT: 01-DEC-2001 (TREMURel, 19, last sequence update)







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068793
ID 068793 PRELIMINARY: PRT: 76 AA.
AC 068793:
DT 01-AUG-1998 (TREMblrel. 07, Created)
DT 01-AUG-1998 (TREMblrel. 07, last sequence update)
DT 01-MAR-2000 (TREMblrel. 20, last annotation update)
DE Hypothetical protein Y1031.
GN Y1031 OR YPM1.600.
OS Yersinia pestis.
OC Plasmid pMT1 (pMT-1).
OC Packer1: Proteobacteria; gamma subdivision; Enterobacteriaceae,
OC Yersinia.
OX NCBI TaxID=632;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=KIM5;
RX MEDLINE=98422474; PubMed=9748454;
RA Hu P., Elliott J., McCready P., Skowronski F., Gaines T.,
RA Kobayashi A., Brubaker R.R., Garcia E.;
RT "Structural organization of virulence-associated plasmids of Yersinia
RT pestis.";
RL J. Bacteriol. 180:5192-5203(1998)
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=KIM10+;
RX MEDLINE=99043899; PubMed=9826348;
RA Lindler T.F., Plano G.V., Buland V., Mayhew G.F., Blattner F.R.;
RT "Complete DNA sequence and detailed analysis of the Yersinia pestis
RT KIM5 plasmid encoding murine toxin and capsular antigen.";
RL Infect. Immun. 66:5731-5742(1998).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=CO-92 / BIVAP ORIENTALIS; PLASMID=PMT1 (PMT 1);
RX MEDLINE=21479413; PubMed=11586360;
RA Parkhill J., Wren B.W., Thomson N.P., Titchall P.W., Holden M.T.G.,
RA Prentice M.B., Sebatilla M., James F.P., Chichester C., Mungall K.L.,
RA Baker S., Basham P., Bentley S.D., Brooks K., Cerdano-Tarraga A.M.,
RA Chillingworth T., Cronin A., Davies P.M., Davis P., Dougan G.,
RA Fellwell T., Hamlin N., Harty S., Japels F., Kutyshov A.V.,
RA Leather S., Mouton S., Oyston P.C.F., Quail M., Rutherford K.,
RA Simmonds M., Skellern J., Stevens K., Whitbread S., Bartell R.G.;
RT "Genome sequence of Yersinia pestis, the causative agent of plague",
RL Nature 413:523-527(2001).
DR EMBL; AF053947; AAC13243.1;
DR EMBL; AF074611; AAC82749.1;
DR EMBL; AL117211; CAB55242.1;
KW Hypothetical protein; Plasmid; Complete proteome.
SQ
SEQUENCE 76 AA: 8407 MW: 78449774506PB 07664;

Query March 100.0%; Score 31; PP 1%; Length 76;
Best Local Similarity 50.0%; Pred. No. 7 rev02;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 50 WRTWNP 55

RESULT 14
O9S4W3 PRELIMINARY: PRT: 77 AA.
AC O9S4W3:
DT 01-MAY-2000 (TREMblrel. 13, Created)
DT 01-MAY-2000 (TREMblrel. 13, last sequence update)
DT 01-MAY-2000 (TREMblrel. 13, last annotation update)
DE Hypothetical 9.1 kDa protein.
OS Escherichia coli.
OC Plasmid P.
OC Packer1: Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.

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FX MEDLINE=99266678; PubMed=10766527;
RA Manwaring N.P., Skurray R.A., Firth N.;
RT "Nucleotide sequence of the P plasmid leading region.";
PL Plasmid 41:219-225(1999).
DR EMBL; AF106329; AAC47189.1;
KW Hypothetical protein; plasmid.
SQ
SEQUENCE 77 AA: 9089 MW: 384345602/85776 28264;

Query March 100.0%; Score 31; PP 2%; Length 77;
Best Local Similarity 50.0%; Pred. No. 7 rev02;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 53 WRTWNP 58

RESULT 15
O9WTF7 PRELIMINARY: PRT: 79 AA.
AC O9WTF7:
DT 01-NOV-1999 (TREMblrel. 12, Created)
DT 01-NOV-1999 (TREMblrel. 12, last sequence update)
DT 01-DEC-2001 (TREMblrel. 19, last annotation update)
DE Ydea protein.
GN YDEA.
OS Escherichia coli.
OC Plasmid p100.
OC Packer1: Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.
RA Sampei G., Mizobuchi K.;
RT "Organization and diversification of plasmid genomes: complete
RT nucleotide sequence of the p100 genome.";
PI Submitted (MAY 1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB000342; BAA78827.1;
KW SEQUENCE 79 AA: 3308 MW: 717648560/1566 7374;

Query March 100.0%; Score 31; PP 2%; Length 79;
Best Local Similarity 50.0%; Pred. No. 7 rev02;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 55 WRTWNP 60

Search completed February 20, 2003, 10:42:20
Job time: 05.333 secs

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GenCore version 5.1.3  
Copyright (c) 1991 - 2003 CompuGen Ltd.

OM protein protein search, using sw method

Run on: February 20, 2003, 11:21:07, Search time 29.333 seconds  
(without alignments)  
27.256 Million cell updates/sec

Title: US-09-912-414-2

Perfect score: 6

Sequence: 1 WVPWHP 6

Scoring table:

Gapop 60 0, Gapext 60 0

Searched: 908470 seqs, 13256620 positions

Word size: 0

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 similarities

Database: A.Geneseq.101002.\*

```
1: /SID2/gcgdata/geneseq/geneexp-emb1/AA1940.DAT.*
2: /SID2/gcgdata/geneseq/geneexp-emb1/AA1941.DAT.*
3: /SID2/gcgdata/geneseq/geneexp-emb1/AA1942.DAT.*
4: /SID2/gcgdata/geneseq/geneexp-emb1/AA1943.DAT.*
5: /SID2/gcgdata/geneseq/geneexp-emb1/AA1944.DAT.*
6: /SID2/gcgdata/geneseq/geneexp-emb1/AA1945.DAT.*
7: /SID2/gcgdata/geneseq/geneexp-emb1/AA1946.DAT.*
8: /SID2/gcgdata/geneseq/geneexp-emb1/AA1947.DAT.*
9: /SID2/gcgdata/geneseq/geneexp-emb1/AA1948.DAT.*
10: /SID2/gcgdata/geneseq/geneexp-emb1/AA1949.DAT.*
11: /SID2/gcgdata/geneseq/geneexp-emb1/AA1950.DAT.*
12: /SID2/gcgdata/geneseq/geneexp-emb1/AA1951.DAT.*
13: /SID2/gcgdata/geneseq/geneexp-emb1/AA1952.DAT.*
14: /SID2/gcgdata/geneseq/geneexp-emb1/AA1953.DAT.*
15: /SID2/gcgdata/geneseq/geneexp-emb1/AA1954.DAT.*
16: /SID2/gcgdata/geneseq/geneexp-emb1/AA1955.DAT.*
17: /SID2/gcgdata/geneseq/geneexp-emb1/AA1956.DAT.*
18: /SID2/gcgdata/geneseq/geneexp-emb1/AA1957.DAT.*
19: /SID2/gcgdata/geneseq/geneexp-emb1/AA1958.DAT.*
20: /SID2/gcgdata/geneseq/geneexp-emb1/AA1959.DAT.*
21: /SID2/gcgdata/geneseq/geneexp-emb1/AA1960.DAT.*
22: /SID2/gcgdata/geneseq/geneexp-emb1/AA1961.DAT.*
23: /SID2/gcgdata/geneseq/geneexp-emb1/AA1962.DAT.*
```

Pred No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Openy Match length	ID	Description
1	6	100.0	6 21 AAB01492	peptide which bind
2	5	83.3	6 21 AAB01504	peptide which bind
3	5	83.3	6 21 AAB01509	peptide which bind
4	5	83.3	372 20 AAY09882	Human LPA receptor
5	5	83.3	373 20 AAY05493	Human RSP 24 prote
6	4	66.7	373 22 AAU00227	LPA receptor-relat
7	4	66.7	6 14 AAR37389	peptide for treati
8	4	66.7	6 21 AAB01505	peptide which bind
9	4	66.7	6 21 AAB01508	peptide which bind
10	4	66.7	10 17 AAR86140	Anti-ELAM-1 bindin

11	4	66.7	10 17 AAR86145	Anti-ELAM-1 bindin
12	4	66.7	10 17 AAR86146	Anti-ELAM-1 bindin
13	4	66.7	10 19 AAM63963	ELAM-1 peptide mim
14	4	66.7	10 19 AAM63964	ELAM-1 peptide mim
15	4	66.7	10 19 AAM63958	ELAM-1 peptide mim
16	4	66.7	12 14 AAR36519	ELAM-1 peptide mim
17	4	66.7	12 15 AAR66756	Random peptide #43
18	4	66.7	12 17 AAR31534	ELAM-1 functional
19	4	66.7	12 18 AAR65286	Anti-body R32.39 op
20	4	66.7	12 18 AAM80380	Peptide eluted aft
21	4	66.7	14 23 AAR74383	Peptide eluted aft
22	4	66.7	27 22 AAR12264	Peptide eluted aft
23	4	66.7	31 22 AAR843615	Peptide #11121 enc
24	4	66.7	31 22 ABB30569	Protein #8766 enc
25	4	66.7	31 22 AAM64556	Human brain cytos
26	4	66.7	31 22 AAM77350	Human bone marrow
27	4	66.7	31 22 AAM21282	Peptide #7716 enc
28	4	66.7	31 22 AAM37509	Peptide #11946 enc
29	4	66.7	31 22 ABB46373	Human peptide enc
30	4	66.7	35 22 ABB47677	Human peptide enc
31	4	66.7	37 22 AAB13641	Human polypeptide
32	4	66.7	38 22 ABB30966	Peptide #4617 enc
33	4	66.7	38 22 ABB36158	Peptide #4617 enc
34	4	66.7	38 22 ABB21544	Protein #1543 enc
35	4	66.7	38 22 AAM69317	Human bone marrow
36	4	66.7	38 22 AAB12011	Human polypeptide
37	4	66.7	38 22 AAM17157	Peptide #1591 enc
38	4	66.7	38 22 AAM29648	Peptide #5685 enc
39	4	66.7	38 22 AAM04851	Peptide #4533 enc
40	4	66.7	38 22 ABB38943	Human peptide enc
41	4	66.7	41 22 AAM13103	Human polypeptide
42	4	66.7	42 22 AAM73279	Human bone marrow
43	4	66.7	42 22 AAM13855	Peptide #1502 enc
44	4	66.7	48 22 ABB64316	Drosophila melanog
45	4	66.7	48 22 AAB11862	Human polypeptide

#### ALIGNMENTS

RESULT 1	AA01492	standard, peptide, 6 AA.
ID	AA01492	
XX	AA01492	
XX	09 NOV 2000 (first entry)	
DT	09 NOV 2000	
DE	Peptide which binds to transcription factor ECF: DNA binding domain.	
XX		
XX	DNA binding; transcription factor; ECF, ECF-1, -2, -3, -4, -5, -6, -7, -8, -9, -10, -11, -12, -13, -14, -15, -16, -17, -18, -19, -20, -21, -22, -23, -24, -25, -26, -27, -28, -29, -30, -31, -32, -33, -34, -35, -36, -37, -38, -39, -40, -41, -42, -43, -44, -45, -46, -47, -48, -49, -50, -51, -52, -53, -54, -55, -56, -57, -58, -59, -60, -61, -62, -63, -64, -65, -66, -67, -68, -69, -70, -71, -72, -73, -74, -75, -76, -77, -78, -79, -80, -81, -82, -83, -84, -85, -86, -87, -88, -89, -90, -91, -92, -93, -94, -95, -96, -97, -98, -99, -100, -101, -102, -103, -104, -105, -106, -107, -108, -109, -110, -111, -112, -113, -114, -115, -116, -117, -118, -119, -120, -121, -122, -123, -124, -125, -126, -127, -128, -129, -130, -131, -132, -133, -134, -135, -136, -137, -138, -139, -140, -141, -142, -143, -144, -145, -146, -147, -148, -149, -150, -151, -152, -153, -154, -155, -156, -157, -158, -159, -160, -161, -162, -163, -164, -165, -166, -167, -168, -169, -170, -171, -172, -173, -174, -175, -176, -177, 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KW human cell invasion, neurotransmitter release, chemotaxis, cancer;  
 KW drug design; tumour; anti coagulant; vascular restenosis, arthritis;  
 KW anti-inflammatory; wound healing; tissue regeneration; blood coagulation;  
 KW nerve regeneration; osteoporosis; bone regeneration; cosmetic; scarring;  
 KW ischaemic heart disease; myocardial infarction; angina pectoris; stroke;  
 KW cerebral infarction; cerebral sequelae; subarachnoid haemorrhage;  
 KW benign prostatic hyperplasia; diabetes mellitus; retinopathy; neuropathy;  
 KW neuropathy; diabetic retinopathy;  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO9929887-A1.  
 XX  
 PD 17-JUN-1999.  
 XX  
 PF 10-DEC-1998; 98WO-US26262.  
 XX  
 PR 10-FEB-1997; 97US-0987943.  
 XX  
 PA (SYNA-) SYNAPTIC PHARM CORP.  
 XX  
 PI Bard JA;  
 XX  
 DR WPI: 1999-385620/32.  
 DR N-PSDB; AAX74130.  
 XX  
 PT Lysophosphatidic acid receptor sequence  
 PS Claim 9; Fig 3A-B, 12pp; English.  
 XX  
 CC This invention describes a novel nucleic acid (1) encoding a human  
 CC lysophosphatidic acid (LPA; 1-acyl-glycerol-3-phosphate) receptor. Probes  
 CC and antibodies which bind to the LPA receptor can be used for detecting  
 CC expression of a mammalian LPA receptor. The probe can be used for  
 CC detecting a predisposition to a disorder associated with LPA receptor  
 CC activity. The transgenic animals incorporating the LPA receptor can be  
 CC used for determining the physiological effects of varying levels of  
 CC activity of mammalian LPA receptors, or for identifying an (ant)agonist  
 CC capable of alleviating a physical or behavioural abnormality associated  
 CC with activity of LPA receptor. LPA receptors produce a wide range of  
 CC responses such as cell proliferation, stimulation of neurite retraction,  
 CC platelet aggregation, smooth muscle contraction, tumour cell invasion,  
 CC neurotransmitter release, chloride efflux and chemotaxis. The human LPA  
 CC receptor can serve as a tool for the drug design of novel  
 CC therapeutic agents for various indications, including cancer, tumour  
 CC reduction or prevention, platelet aggregation (as an anti-coagulant),  
 CC vascular restenosis, arthritis (as an anti-inflammatory), wound healing,  
 CC tissue regeneration (preferably skin and nerve regeneration), blood  
 CC coagulation, osteoporosis (bone regeneration), and cosmetic uses  
 CC (preferably the prevention of abnormal growths or scarring or for  
 CC augmentation). This receptor can additionally serve as a tool for the  
 CC drug design of novel therapeutic agents for indications including  
 CC ischaemic heart diseases (e.g. myocardial infarction and angina  
 CC pectoris), stroke due to cerebrovascular disturbances (e.g. cerebral  
 CC infarction and cerebral sequelae after subarachnoid haemorrhage), benign  
 CC prostatic hyperplasia, complications of diabetes mellitus (e.g.  
 CC retinopathy, neuropathy, and nephropathy), and diabetic retinopathy.  
 XX  
 SQ Sequence 372 AA;  
 Query Match 83.3%; Score 5, DB 20; Length 372;  
 Best Local Similarity 100.0%; Prod. No. 8;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0.

AC AAY05493;  
 XX  
 DT 07-JUL-1999 (first entry)  
 XX  
 DE Human PSP-24 protein sequence.  
 XX  
 KW FFG-1, FFG-2, FFG-3, FFG 4, FFG 5, PSP 24, human, detecting therapy;  
 KW inverse agonist; allosteric modulation; lysophosphatidic acid receptors;  
 KW LPA signalling mediated disease; cellular apoptosis.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO9919513-A2.  
 XX  
 PD 22-APR-1999.  
 XX  
 PF 09-OCT-1998; 98WO-US21315.  
 XX  
 PR 10-OCT-1997; 97US-0061572.  
 XX  
 PA (LXPB-) LXP BIOTECHNOLOGY INC  
 XX  
 PI Erikson J, Goddard JG, Kiefer M;  
 XX  
 DR WPI: 1999-277658/23.  
 DR N-PSDB; AAX36570.  
 XX  
 PT Identification of (ant)agonists of LPA receptor EDG-2 (or, e.g.,  
 PT treating LPA signalling mediated diseases such as cellular apoptosis)  
 XX  
 PS Claim 1; Page 58-60; 63pp; English.  
 XX  
 CC This sequence represents human PSP-24.  
 CC The invention relates to methods of detecting (ant)agonist, inverse  
 CC agonist or allosteric modulators of the lysophosphatidic acid receptors  
 CC EDG-1, EDG-2, FFG-3, FFG-4, FFG-5, and PSP-24. The methods are used to  
 CC identify (ant)agonists and allosteric modulators of the lysophosphatidic  
 CC acid (LPA) EDG2 receptor, e.g. to treat LPA signalling mediated disease  
 CC such cellular apoptosis.  
 XX  
 SQ Sequence 373 AA;  
 Query Match 83.3%; Score 5, DB 20; Length 373;  
 Best Local Similarity 100.0%; Prod. No. 8;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Cy 2 VRWHF 6  
 Db 98 VRWHF 102  
 XX  
 RESULT 6  
 ID AAY05493 standard; Protein: 373 AA.  
 XX  
 AC AAY05493;  
 XX  
 DT 11-MAY-2001 (first entry)  
 XX  
 DE LPA receptor-related amino acid sequence #1.  
 XX  
 KW LPA receptor, EDG-2, lysophosphatidic acid, phospholipid, tumour;  
 KW cell signalling, MAP kinase; LPA modulator; neurodegenerative disease;  
 KW Alzheimer's disease; Parkinson's disease; neuron damage; apoptosis;  
 KW ischaemic heart disease; viral; HIV; inflammatory bowel disease;  
 KW organ transplant; mouse.  
 XX  
 OS Mus sp.  
 XX  
 PN WO200112838-A2.  
 XX  
 PD 22-FEB-2001.  
 XX

XX	11-AUG-1999	99US-0658744.	
XX	18-AUG-1999	99US-0376839.	
XX	(ATAI) AIRBORN TECHNOLOGIES INC		
XX	ENTERPRISE, 243043 02, 90650 M, F1000 02		
XX	MDA, 2001 020507003		
XX	N-PSDB; AAS00260.		
XX	Medical technology of a biotechnology firm is the primary for		
XX	developing vaccines, other diseases, including including the use of		
XX	recombinant DNA technology, producing drugs of value in the treatment of		
XX	diabetes, Lyme disease, HIV, English.		
XX	The company's research, the firm is currently conducting research in		
XX	used during analysis of 2,2,6,6-tetramethyl-4-pyridone (TMP), EDC, the		
XX	reagent is given in the specification but no further information is		
XX	given. This is a highly purified form of a variety of plant and animal		
XX	materials, including plant and animal materials, including a		
XX	Myeloid leukemia, acute (MEL) or B-cell lymphoma (BCL) involves involvement		
XX	a 2,2,6,6-tetramethyl-4-pyridone (TMP) derivative, the use of which the		
XX	reagent is used in the treatment of diseases characterized by slow growth of		
XX	repair of neuronal cells, neurodegenerative diseases, such as Alzheimer's		
XX	disease, Parkinson's disease, and acute neuron damage, for maintaining		
XX	apoptotic pathways and repairing damaged brain tissue, for use in		
XX	diagnosed liver disease, and rejection of organ transplants.		
XX	Sequence 402 AA)		
XX	Query March 81.3%; Score 5; DB 22; Length 373;		
XX	Best Local Similarity 100.0%; E-Exp. No. 9;		
XX	Matches 2; Conservation 0; Mismatches 0; Indels 0; Gaps 0;		
XX	QY 2 VRWHP 6		
XX	DB 98 VRWHP 102		
XX	RESUME 7		
XX	AAR17389		
XX	ID AAR17389 standard; peptide; 6 AA.		
XX	XX AAR17389;		
XX	DT 07-JUN-1993 (first entry)		
XX	DE Peptide for treating with shock.		
XX	XX Toxic shock; blood anti-toxin removal; acute diagnosis; sepsis;		
XX	XX anti-toxin; polypeptide; anti-toxin; polypeptide; bacterial meningitis;		
XX	XX HIV related infections; polymyxin B; Group II.		
XX	XX Synthetic.		
XX	XX Key		
XX	XX Region		
XX	XX Region		
XX	XX Location/Qualifiers		
XX	XX 1		
XX	XX /note "repeat region"		
XX	XX 4..6		
XX	XX /note="repeat region"		
XX	XX ZA9200943-A.		
XX	XX 25-NOV-1992.		
XX	XX 10-FEB-1992;		
XX	XX 92ZA 0000443.		
XX	XX 11-FEB-1992;		
XX	XX 91US-0658744.		

XX (PDB/1-PEP M.  
 XX Folio M;  
 XX WP1; 1991-04/04/11.  
 XX The peptide 61 (treatment of prevention of 1991-04-11) comprises  
 XX specified sequences of amino acids) and analogs  
 XX including sequences related thereto.  
 XX  
 XX Example; Page 5; 39pp; English.  
 XX  
 XX The Group 11 peptide is an example of a protein peptide of formula  
 XX  $R_1 - (Gly)_n - R_2$ , where  $R_1$  is a peptide chain of 1-100  
 XX amino acids,  $R_2$  is a peptide chain of 1-100 amino acids, and  
 XX each  $P$  is H, an amino acid residue or a fatty acid residue.  
 XX The peptide is useful for treating or preventing septic shock,  
 XX related with a disease, P1, P2, P3, P4, P5, P6, P7, P8, P9, P10, P11,  
 XX P12, P13, P14, P15, P16, P17, P18, P19, P20, P21, P22, P23, P24, P25,  
 XX P26, P27, P28, P29, P30, P31, P32, P33, P34, P35, P36, P37, P38, P39,  
 XX P40, P41, P42, P43, P44, P45, P46, P47, P48, P49, P50, P51, P52, P53,  
 XX P54, P55, P56, P57, P58, P59, P60, P61, P62, P63, P64, P65, P66, P67,  
 XX P68, P69, P70, P71, P72, P73, P74, P75, P76, P77, P78, P79, P80, P81,  
 XX P82, P83, P84, P85, P86, P87, P88, P89, P90, P91, P92, P93, P94, P95,  
 XX P96, P97, P98, P99, P100, P101, P102, P103, P104, P105, P106, P107,  
 XX P108, P109, P110, P111, P112, P113, P114, P115, P116, P117, P118, P119,  
 XX P120, P121, P122, P123, P124, P125, P126, P127, P128, P129, P130, P131,  
 XX P132, P133, P134, P135, P136, P137, P138, P139, P140, P141, P142, P143,  
 XX P144, P145, P146, P147, P148, P149, P150, P151, P152, P153, P154, P155,  
 XX P156, P157, P158, P159, P160, P161, P162, P163, P164, P165, P166, P167,  
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 XX P180, P181, P182, P183, P184, P185, P186, P187, P188, P189, P190, P191,  
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 XX P228, P229, P230, P231, P232, P233, P234, P235, P236, P237, P238, P239,  
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 XX P720, P721, P722, P723, P724, P725, P726, P727

```

XX XX Example; Page 26; 42pp; English.
PS
CC CC Peptides which bind to the DNA binding domain of transcription
CC factor E2F and inhibit cell cycle progression may be useful as
CC research agents to investigate the interaction between E2F and DP-1,
CC or the activation of transcription by E2F-1/DP-1 heterodimers. They
CC may also be used for inducing apoptosis and/or cell cycle arrest in
CC a cell, particularly for treatment of cancer or other proliferative
CC disorders such as psoriasis and restenosis.
XX
XX Sequence 6 AA:
SO
Query Match 66.7%; Score 4; DB 21; Length 6;
Best Local Similarity 100.0%; Pred No. 7 seqs
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0
CY 3 RWHP 6
   |||
   |||
Dh 3 RWHP 6
RESULT 9
ID AAB01508 standard; peptide; 6 AA.
AAB01508
XX AAB01508;
XX
XX 0A NOV 2000 (first entry)
XX
DE Peptide which binds to transcription factor E2F-1 DNA binding domain
XX
XX DNA binding; transcription factor; E2F, E2F 1, cell cycle, DP 1,
XX activation; transcription; apoptosis; proliferative disorder;
XX psoriasis; restenosis.
XX
XX Synthetic.
XX
XX WO200044771-A1.
XX
XX 03-AUG-2000.
XX
XX 26-JAN 2000; 266AWP (P00027).
XX
XX 26-JAN-1999; 94RF-0001710.
XX
XX (PROL-1) PROLIFIX LTD.
XX
XX Mueller P, Kontermann PE, Montigiani S;
XX
XX WPI: 2000-532806/48.
XX
XX Peptides binding to the DNA binding domain of transcription factor E2F
XX and inhibiting cell cycle progression, useful for the treatment of
XX cancer
XX
XX Example; Page 26; 42pp; English
XX
XX Peptides which bind to the DNA binding domain of transcription
XX factor E2F and inhibit cell cycle progression may be useful as
XX research agents to investigate the interaction between E2F and DP-1,
XX or the activation of transcription by E2F-1/DP-1 heterodimers. They
XX may also be used for inducing apoptosis and/or cell cycle arrest in
XX a cell, particularly for treatment of cancer or other proliferative
XX disorders such as psoriasis and restenosis.
XX
XX Sequence 6 AA:
SO
Query Match 66.7%; Score 4; DB 21; Length 6;
Best Local Similarity 100.0%; Pred No. 7 seqs;
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0
CY 1 WVRW 4

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Db      1 WVRW 4
|||||
RESULT 10
AAAR86140
1D  AAR86140 standard, ref136, 10 AA.
XX
XX
AC  AAR86140:
XX
DT  26-JUN-1996 (first entry)
XX
DE  Anti-ELAM-1 binding peptide #117.
XX
XX  Peptide #117:117, endothelial leukocyte adhesion molecule, ELAM, selectin;
XX  receptor, leukocyte, vascular wall, endothelium, extravasation;
XX  inflammation; steryl Lewisx cell surface glycoprotein; HMO cells.
XX
XX  Synthetic.
XX
XX  WO9531210-A1.
XX
XX  23-NOV-1995.
XX
XX  11 MAY 1995, 95WO-US06315.
XX
XX  11-MAY-1994, 94US-0241054.
XX
XX  (AFTV-) AFTVMAX TECHNOLOGIES INC.
XX
XX  Barrett BW, Twirla SE, Dower WJ, Koller KL, Lee J;
XX  Martens CL, Ruhland-Fritsch B;
XX  WPI, 1996-010607/01.
XX
XX  New peptide(s) that bind to endothelial leukocyte adhesion molecule
XX  1 - useful for treating inflammation and other E-selectin mediated
XX  diseases
XX
XX  Disclosure; Page 17; 85pp; English.
XX
XX  Peptides AAR86124-AAR86236 are examples of peptides and their mimics
XX  that bind to endothelial leukocyte adhesion molecule (ELAM-1) and
XX  molecule is a member of the selectin family of receptors and is involved
XX  in binding of leukocytes to the vascular endothelial wall prior to
XX  extravasation of the leukocyte, e.g. to a site of inflammation.
XX  The peptides bind prefer to E-selectin but may also bind to or
XX  E-selectin, and can be used to treat conditions mediated by E-selectin,
XX  e.g. inflammatory conditions. The peptides have strong affinity for the
XX  selectin receptors and inhibit the binding of the steryl Lewisx (also x)
XX  part of cell surface glycoproteins to E-selectin. The peptide are
XX  small, generally less than 20, have an  $\alpha$  of 10 to 100 amino acids
XX  adjacent binding of HMO cells to ELAM-1, have one or more peptide
XX  linkages replaced by CH2CH2CH2CH2 phosphate, CH2CH2CH2CH2CONH2,
XX  or NHCOOH linkages where R = H or a lower alkyl and Pd a lower alkyl.
XX  The peptides may also have substituted N- and C-termini e.g.
XX  succinyl, N-benzyloxycarbonyl or a lower alkyl cids.
XX
XX  Sequence 10 AA:
XX
XX  Query Match 66.7% Score 4; FB 17; Length 10;
XX  Fast Local Similarity 100.0%, fixed. No. 9.6;
XX  Matches 4, Conservation 0, Widespread 0; Update 0; Opt 0;
XX
XX  1 WVRW 4
XX  |||||
XX  DB 6 WVRW 9
XX
XX  RESULT 11
XX  AAR86145
XX  1D  AAR86145 standard; peptide; 10 AA.
XX

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FT Modified-site 10 /note="C-terminal Met is amidated"  
 FT XX  
 PN US5728802-A.  
 XX  
 PD 17-MAR-1998.  
 XX  
 PF 12-MAY-1995; 92US-0439817.  
 XX  
 PR 12-MAY-1995; 95US-0439817.  
 PR 06-MAY-1992; 92US-0881395.  
 PR 05-MAY-1993; 93US-0057295.  
 PR 11-MAY-1994; 94US-0241054.  
 XX  
 PA (AFFY-) AFFYMAX TECHNOLOGIES NV.  
 XX  
 PI Barrett RW, Cwirla SE, Dower WJ, Koller KJ, Lee J;  
 PI Martens CL, Ruhland B;  
 XX  
 DP WPI; 1998-249882/22.  
 XX  
 PT Peptide(s) or their mimetic(s) that bind to E-selectin - useful for,  
 PT e.g. treating conditions mediated by E-selectin such as inflammatory  
 PT condition(s)  
 XX  
 PS Example 2; Column 93-94; 84pp; English.  
 XX  
 CC AAM63846-W64054 are peptides and peptide mimetics that bind selecting  
 CC including endothelial leukocyte adhesion molecule 1 (ELAM-1) and can be  
 CC used for blocking adhesion of leukocytes to the selectins. The peptides  
 CC have applications for the treatment of conditions mediated by  
 CC E-selectin, e.g. inflammatory conditions. They can also be used for  
 CC diagnostic purposes, e.g. for identifying the vascular site of E-selectin  
 CC in vivo or can be coupled to anti-inflammatory or other drugs.  
 XX  
 SO Sequence 10 AA;  
 Query Match 66.7%; Score 4; DR 19; Length 10;  
 Best Local Similarity 100.0%; Pred No 9.6;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 WVRW 4  
 DB 6 WVRW 9  
 RESULT 14  
 AAM63964  
 ID AAM63964 standard; peptide; 10 AA.  
 XX  
 AC AAM63964;  
 XX  
 DT 02-OCT-1998 (first entry)  
 XX  
 DE ELAM-1 peptide mimetic #119.  
 XX  
 KW Endothelial leukocyte adhesion molecule 1; ELAM-1; inflammation;  
 KW selectin; diagnosis; mimetic.  
 XX  
 OS Synthetic.  
 XX  
 PN US5728802-A.  
 XX  
 PD 17-MAR-1998.  
 XX  
 PF 12-MAY-1995; 92US-0439817.  
 XX  
 PR 12-MAY-1995; 95US-0439817.  
 PR 06-MAY-1992; 92US-0881395.  
 PR 05-MAY-1993; 93US-0057295.  
 PR 11-MAY-1994; 94US-0241054.  
 XX  
 PA (AFFY-) AFFYMAX TECHNOLOGIES NV.

XX  
 PI Barrett RW, Cwirla SE, Dower WJ, Koller KJ, Lee J;  
 PI Martens CL, Ruhland B;  
 XX  
 DP WPI; 1998-249882/22.  
 XX  
 PT Peptide(s) or their mimetic(s) that bind to E-selectin - useful for,  
 PT e.g. treating conditions mediated by E-selectin such as inflammatory  
 PT condition(s)  
 XX  
 PS Example 2; Column 93-94; 84pp; English.  
 XX  
 CC AAM63846-W64054 are peptides and peptide mimetics that bind selecting  
 CC including endothelial leukocyte adhesion molecule 1 (ELAM-1) and can be  
 CC used for blocking adhesion of leukocytes to the selectins. The peptides  
 CC have applications for the treatment of conditions mediated by  
 CC E-selectin, e.g. inflammatory conditions. They can also be used for  
 CC diagnostic purposes, e.g. for identifying the vascular site of E-selectin  
 CC in vivo or can be coupled to anti-inflammatory or other drugs.  
 XX  
 SO Sequence 10 AA;  
 Query Match 66.7%; Score 4; DR 19; Length 10;  
 Best Local Similarity 100.0%; Pred No 9.6;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 WVRW 4  
 DB 6 WVRW 9  
 RESULT 15  
 AAM63958  
 ID AAM63958 standard; peptide; 10 AA.  
 XX  
 AC AAM63958;  
 XX  
 DT 02-OCT-1998 (first entry)  
 XX  
 DE ELAM-1 peptide mimetic #113.  
 XX  
 KW Endothelial leukocyte adhesion molecule 1; ELAM-1; inflammation;  
 KW selectin; diagnosis; mimetic.  
 XX  
 OS Synthetic.  
 XX  
 PN US5728802-A.  
 XX  
 PD 17-MAR-1998.  
 XX  
 PF 12-MAY-1995; 92US-0439817.  
 XX  
 PR 12-MAY-1995; 95US-0439817.  
 PR 06-MAY-1992; 92US-0881395.  
 PR 05-MAY-1993; 93US-0057295.  
 PR 11-MAY-1994; 94US-0241054.  
 XX  
 PA (AFFY-) AFFYMAX TECHNOLOGIES NV.  
 XX  
 PI Barrett RW, Cwirla SE, Dower WJ, Koller KJ, Lee J;  
 PI Martens CL, Ruhland B;  
 XX  
 DP WPI; 1998-249882/22.  
 XX  
 PT Peptide(s) or their mimetic(s) that bind to E-selectin - useful for,  
 PT e.g. treating conditions mediated by E-selectin such as inflammatory  
 PT condition(s)  
 XX  
 PS Example 2; Column 91-92; 84pp; English.  
 XX  
 CC AAM63846-W64054 are peptides and peptide mimetics that bind selecting  
 CC including endothelial leukocyte adhesion molecule 1 (ELAM-1) and can be  
 CC used for blocking adhesion of leukocytes to the selectins. The peptides

CC Have applications for the treatment of conditions mediated by  
 CC E-selectin, e.g. inflammatory conditions. They can also be used for  
 CC diagnostic purposes, e.g. for identifying the vascular site of E-selectin  
 CC in vivo or can be coupled to anti-inflammatory or other drugs.  
 XX

SO Sequence 10 AA;

Query Match 66.7%; Score 4; DB 19; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 9.6;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRW 4  
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 Db 6 WVRW 9

Search completed: February 20, 2003, 11:26:31  
 Job time: 30.333 secs

GenCore version 5.1.3  
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OM Protein - Protein search, using SW model

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(without alignments)  
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Title: US-09-912 414 2

Perform: Score

Sequence: 1 WVRMHF 6

Search: Filter

Gapop: 60.0, Gapext: 60.0

Search: 468190 seqs, 99997795 residues

Word size: 0

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Post-processing: Listing first 45 summaries

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Pred No is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	83.3	372	6	US-10-057-498-1394	Sequence 394, App
2	66.7	20	1	PCT-US02-34021-373	Sequence 373, App
3	66.7	20	1	PCT-US02-34021-422	Sequence 422, App
4	66.7	38	6	US-10-057-498-13934	Sequence 13934, App
5	66.7	52	1	PCT-US02-32727-5526	Sequence 5526, App
6	66.7	52	1	US-09-978-825-15052	Sequence 15052, App
7	66.7	52	1	US-10-057-498-15052	Sequence 15052, App
8	66.7	52	1	PCT-US02-32727-7189	Sequence 7189, App
9	66.7	56	1	US-09-978-825-7189	Sequence 7189, App
10	66.7	58	1	US-10-057-498-11606	Sequence 11606, App
11	66.7	58	1	PCT-US02-32727-11606	Sequence 11606, App
12	66.7	59	1	US-09-978-825-11606	Sequence 11606, App
13	66.7	59	1	US-10-057-498-11606	Sequence 11606, App
14	66.7	64	1	PCT-US02-32727-44772	Sequence 44772, App
15	66.7	64	1	US-09-978-825-44772	Sequence 44772, App
16	66.7	64	1	US-10-057-498-24772	Sequence 24772, App
17	66.7	65	1	PCT-US02-32727-24493	Sequence 24493, App
18	66.7	65	1	US-09-978-825-24493	Sequence 24493, App
19	66.7	65	1	US-10-057-498-24493	Sequence 24493, App
20	66.7	65	1	US-09-978-825-24493	Sequence 24493, App
21	66.7	66	1	PCT-US02-32727-25628	Sequence 25628, App
22	66.7	66	1	US-09-978-825-25628	Sequence 25628, App
23	66.7	67	1	US-10-057-498-25628	Sequence 25628, App
24	66.7	67	1	US-09-978-825-25628	Sequence 25628, App
25	66.7	70	1	PCT-US02-32727-23166	Sequence 23166, App
26	66.7	70	1	US-09-978-825-23166	Sequence 23166, App

#### ALIGNMENTS

```

RESULT 1
US-10-057-498-1394
Sequence 394, Application US/10225567A
GENERAL INFORMATION:
APPLICANT: Lifespan Biosciences
APPLICANT: Brown, Joseph P.
APPLICANT: Burner, Glenn C.
APPLICANT: Koush, Christine L.
TITLE OF INVENTION: ANTICIPICITILIN AND ANTICIPICITILIN
FILE REFERENCE: 1990-4-4
CURRENT APPLICATION NUMBER: US/10-057-498-1394
CURRENT FILING DATE: 2002-10-19
PRIORITY FILING DATE: 2000-12-19
NUMBER OF SEQ ID NOS: 2092
SOFTWARE: Parent In version 3.1
SEQ ID NO: 394
LENGTH: 372
TYPE: PRT
ORGANISM: Homo sapiens
US-10-057-498-1394

Query Match      83.3% Score 57 DB 6: Length 372:
Post local similarity 100% Pct 53:
Matches 5, Conservative 0, Mismatches 0, Gaps 0

Cy      2 WVRMHF 6
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Db      98 WVRMHF 102

RESULT 2
PCT-US02-34021-373
Sequence 373, Application PC/TUS0234021
GENERAL INFORMATION:
APPLICANT: DCR Biotechnology, Inc.
TITLE OF INVENTION: TAPD57-88-01-010-3-REFINING AND ITS USE IN IDENTIFYING AND SETTING
FILE REFERENCE: 2598-4009PC
CURRENT APPLICATION NUMBER: PCT/US-02/34021
CURRENT FILING DATE: 2002-10-24
PRIORITY FILING DATE: 2001-10-24
NUMBER OF SEQ ID NOS: 537
SOFTWARE: Parent In version 3.1
SEQ ID NO: 373
LENGTH: 20
TYPE: PRT
ORGANISM: Escherichia coli
FEATURE:

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NAME/KEY: MISC FEATURE  
 OTHER INFORMATION: VERRA-44F 4 M  
 PCT-US02-34021-427

Query Match 66.7% Score 4, DB 1, Length 20,  
 Best Local Similarity 100.0%, Pred. No. 88;  
 Matches 4, Conservative 0, Mismatches 0, Indels 0, Gaps 0,

2 YEAR 5  
 11 VPMH 14

Result 4  
 PCT-US02-34021-427  
 Sequence 422, Application: F07H03/34021  
 GENERAL INFORMATION:  
 APPLICANT: DGI Biotechnology, Inc.  
 TITLE OF INVENTION: TARGET SPECIFIC SCREENING AND ITS USE FOR IDENTIFYING TARGET BIND  
 FILE REFERENCE: 2598-4009PC  
 CURRENT APPLICATION NUMBER: PCT/US02/34021  
 PRIOR FILING DATE: 2002-10-24  
 PRIOR FILING DATE: 2001-10-24  
 NUMBER OF SEQ ID NOS: 537  
 SOFTWARE: Patent In Version 3.1  
 SEQ ID NO: 422  
 LENGTH: 20  
 TYPE: PRT  
 ORGANISM: Escherichia coli  
 FEATURE:  
 NAME/KEY: MISC FEATURE  
 OTHER INFORMATION: VERRA-44F-4112  
 PCT-US02-34021-422

Query Match 66.7% Score 4, DB 1, Length 20,  
 Best Local Similarity 100.0%, Pred. No. 88;  
 Matches 4, Conservative 0, Mismatches 0, Indels 0, Gaps 0,

2 YEAR 5  
 11 VPMH 14

Result 4  
 US-10-203-139A-139A  
 Sequence 139A, Application: US/10003139A  
 GENERAL INFORMATION:  
 APPLICANT: Molecular Dynamics, Inc.  
 APPLICANT: Penn, Sharon G  
 APPLICANT: Saito, Toshiro  
 APPLICANT: Hanzel, David K.  
 APPLICANT: Chen, Wensheng  
 TITLE OF INVENTION: HUMAN GENE DERIVED SINGLE EXON NUCLEIC ACID POLYMER COMPOUND FOR  
 TITLE OF INVENTION: ANALOGUE OF GENE EXPRESSION IN BT 474  
 FILE REFERENCE: PB 6004 WO 8  
 CURRENT APPLICATION NUMBER: US/10/203-139A  
 PRIOR FILING DATE: 2002-08-02  
 PRIOR APPLICATION NUMBER: US 60/160,312  
 PRIOR FILING DATE: 24 February 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 26 May 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 03 August 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 03 October 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 17 September 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 21 September 2000  
 PRIOR APPLICATION NUMBER: 00/000,000  
 PRIOR FILING DATE: 10 June 2000  
 NUMBER OF SEQ ID NOS: 15438

SOFTWARE: Molecular Dynamics Oligo 4.1.10.1  
 OTHER INFORMATION: VERRA-44F 4 M  
 PCT-US02-34021-427

Query Match 66.7% Score 4, DB 6, Length 38,  
 Best Local Similarity 100.0%, Pred. No. 14902;  
 Matches 4, Conservative 0, Mismatches 0, Indels 0, Gaps 0,

2 YEAR 5  
 11 VPMH 14

Result 5  
 PCT-US02-32727-5526  
 Sequence 5526, Application: F07H03/32727  
 GENERAL INFORMATION:  
 APPLICANT: Mitcham, Jennifer  
 APPLICANT: Shelly, Yashir  
 APPLICANT: Pelsley, David  
 APPLICANT: Bharat, Ajay  
 APPLICANT: Malhotra, Jeet, Flavio  
 APPLICANT: Zhang, Yanni  
 APPLICANT: Wang, Shiyang  
 APPLICANT: Jen, Shiyang  
 APPLICANT: Lukes, Michael  
 APPLICANT: Benson, Martin  
 APPLICANT: Jones, Robert  
 APPLICANT: Galt, David  
 APPLICANT: Barth, Brenda  
 APPLICANT: Tardiff, John  
 TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acne W  
 FILE REFERENCE: 21021, 51401  
 CURRENT APPLICATION NUMBER: PCT/US02/32727  
 CURRENT FILING DATE: 2002-10-11  
 NUMBER OF SEQ ID NOS: 30992  
 SEQ ID NO: 5526  
 LENGTH: 52  
 TYPE: PRT  
 ORGANISM: Propionibacterium  
 PCT-US02-32727-5526

Query Match 66.7% Score 4, DB 1, Length 52,  
 Best Local Similarity 100.0%, Pred. No. 14902;  
 Matches 4, Conservative 0, Mismatches 0, Indels 0, Gaps 0,

2 YEAR 5  
 26 VPMH 29

Result 6  
 US-10-203-139A-139A  
 Sequence 139A, Application: US/10003139A  
 GENERAL INFORMATION:  
 APPLICANT: Mitcham, Jennifer  
 APPLICANT: Shelly, Yashir  
 APPLICANT: Pelsley, David  
 APPLICANT: Bharat, Ajay  
 APPLICANT: Malhotra, Jeet, Flavio  
 APPLICANT: Zhang, Yanni  
 APPLICANT: Wang, Shiyang  
 APPLICANT: Jen, Shiyang  
 APPLICANT: Lukes, Michael  
 APPLICANT: Benson, Martin  
 APPLICANT: Jones, Robert



```

; APPLICANT: Carter, Derrick
; APPLICANT: Barth, Brenda
; APPLICANT: Douglas, John
; TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acnes V
; FILE REFERENCE: 210121.51401
; CURRENT APPLICATION NUMBER: US/09/978,825
; CURRENT FILING DATE: 2003-01-20
; NUMBER OF SEQ ID NOS: 10992
; SEQ ID NO: 5526
; LENGTH: 52
; TYPE: PRT
; ORGANISM: Propionibacterium
US-09-978-825-5526

Query Match
Best Local Similarity: 100.0%; Score 4; DB 5; Length 52;
Matches: 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5
Db 26 VRMH 29

RESULT 7
US-10-057-498-5526
; Sequence 5526, Application US/10057498
; GENERAL INFORMATION:
; APPLICANT: Mitcham, Jennifer
; APPLICANT: Skeiky, Yasir
; APPLICANT: Persing, David
; TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acnes V
; FILE REFERENCE: 210121.514
; CURRENT APPLICATION NUMBER: US/10/057,499
; CURRENT FILING DATE: 2003-04-20
; NUMBER OF SEQ ID NOS: 29212
; SEQ ID NO: 5526
; LENGTH: 52
; TYPE: PRT
; ORGANISM: Propionibacterium
US-10-057-498-5526

Query Match
Best Local Similarity: 66.7%; Score 4; DB 6; Length 52;
Matches: 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5
Db 26 VRMH 29

RESULT 8
PCT-US02-32727-7189
; Sequence 7189, Application PCT/US0232727
; GENERAL INFORMATION:
; APPLICANT: Mitcham, Jennifer
; APPLICANT: Skeiky, Yasir
; APPLICANT: Persing, David
; APPLICANT: Bhatia, Ajay
; APPLICANT: Malsomneuve, Jean Francois
; APPLICANT: Zhang, Yanni
; APPLICANT: Wang, Siqing
; APPLICANT: Jen, Shiyuan
; APPLICANT: Lodes, Michael
; APPLICANT: Benson, Darin
; APPLICANT: Jones, Robert
; APPLICANT: Carter, Derrick
; APPLICANT: Barth, Brenda
; APPLICANT: Douglas, John
; TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acnes V
; FILE REFERENCE: 210121.51401
; CURRENT APPLICATION NUMBER: PCT/US02/32727
; CURRENT FILING DATE: 2002-10-11
; NUMBER OF SEQ ID NOS: 30992

```

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; SEQ ID NO 7189
; LENGTH: 56
; TYPE: PRT
; ORGANISM: Propionibacterium
PCT-US02-32727-7189

Query Match
Best Local Similarity: 66.7%; Score 4; DB 5; Length 56;
Matches: 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5
Db 20 VRMH 23

RESULT 9
US-09-978-825-7189
; Sequence 7189, Application US/09978825
; GENERAL INFORMATION:
; APPLICANT: Mitcham, Jennifer
; APPLICANT: Skeiky, Yasir
; APPLICANT: Persing, David
; APPLICANT: Bhatia, Ajay
; APPLICANT: Malsomneuve, Jean Francois
; APPLICANT: Zhang, Yanni
; APPLICANT: Wang, Siqing
; APPLICANT: Jen, Shiyuan
; APPLICANT: Lodes, Michael
; APPLICANT: Benson, Darin
; APPLICANT: Jones, Robert
; APPLICANT: Carter, Derrick
; APPLICANT: Barth, Brenda
; APPLICANT: Douglas, John
; TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acnes V
; FILE REFERENCE: 210121.51401
; CURRENT APPLICATION NUMBER: US/09/978,825
; CURRENT FILING DATE: 2003-01-20
; NUMBER OF SEQ ID NOS: 30992
; SEQ ID NO 7189
; LENGTH: 56
; TYPE: PRT
; ORGANISM: Propionibacterium
US-09-978-825-7189

Query Match
Best Local Similarity: 66.7%; Score 4; DB 5; Length 56;
Matches: 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5
Db 20 VRMH 23

RESULT 10
US-10-057-498-7189
; Sequence 7189, Application US/10057498
; GENERAL INFORMATION:
; APPLICANT: Mitcham, Jennifer
; APPLICANT: Skeiky, Yasir
; APPLICANT: Persing, David
; APPLICANT: Bhatia, Ajay
; APPLICANT: Malsomneuve, Jean Francois
; APPLICANT: Zhang, Yanni
; APPLICANT: Wang, Siqing
; APPLICANT: Jen, Shiyuan
; APPLICANT: Lodes, Michael
; APPLICANT: Benson, Darin
; APPLICANT: Jones, Robert
; APPLICANT: Carter, Derrick
; APPLICANT: Barth, Brenda
; APPLICANT: Douglas, John
; TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acnes V
; FILE REFERENCE: 210121.514
; CURRENT APPLICATION NUMBER: US/10/057,498
; CURRENT FILING DATE: 2003-04-20
; NUMBER OF SEQ ID NOS: 30992
; SEQ ID NO 7189
; LENGTH: 56
; TYPE: PRT
; ORGANISM: Propionibacterium
US-10-057-498-7189

Query Match
Best Local Similarity: 66.7%; Score 4; DB 6; Length 56;
Matches: 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5  
| | | |  
DB 20 VRMH 23

RESULT 11  
PCT-US02.32727-11626

Sequence 11626, Application PC/US0232727

GENERAL INFORMATION:  
APPLICANT: Mitcham, Jennifer  
APPLICANT: Skeiky, Yasir  
APPLICANT: Persing, David  
APPLICANT: Bhutia, Ajay  
APPLICANT: Maisonneuve, Jean Francois  
APPLICANT: Zhang, Yanni  
APPLICANT: Jey, Chyian  
APPLICANT: Jey, William  
APPLICANT: Benson, Darin  
APPLICANT: Jones, Robert  
APPLICANT: Carter, Darick  
APPLICANT: Baruch, Brenda  
APPLICANT: Douglas, John  
TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Auto V  
FILE REFERENCE: 210121-51401  
CURRENT APPLICATION NUMBER: PCT/US02/32727  
CURRENT FILING DATE: 2002-10-11  
NUMBER OF SEQ ID NOS: 39992  
SEQ ID NO 11626  
LENGTH: 58  
TYPE: PRT  
ORGANISM: Propionl acnes  
PCT US02.32727-11626

Query Match 66.7%; Score 4; DB 1; Length 58;  
Best Local Similarity 100.0%, Fred. No. 2e+02;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5  
| | | |  
DB 11 VRMH 14

RESULT 12

US 09-978-825-11626

Sequence 11626, Application US/09978825

GENERAL INFORMATION:  
APPLICANT: Mitcham, Jennifer  
APPLICANT: Skeiky, Yasir  
APPLICANT: Persing, David  
APPLICANT: Bhutia, Ajay  
APPLICANT: Maisonneuve, Jean Francois  
APPLICANT: Zhang, Yanni  
APPLICANT: Jey, Chyian  
APPLICANT: Jey, William  
APPLICANT: Benson, Darin  
APPLICANT: Jones, Robert  
APPLICANT: Carter, Darick  
APPLICANT: Baruch, Brenda  
APPLICANT: Douglas, John  
TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Auto V  
FILE REFERENCE: 210121-51401  
CURRENT APPLICATION NUMBER: US/09/978,825  
CURRENT FILING DATE: 2003-01-29  
NUMBER OF SEQ ID NOS: 39992  
SEQ ID NO 11607  
LENGTH: 58  
TYPE: PRT  
ORGANISM: Propionl acnes  
US 09-978-825-11626

Query Match 66.7%; Score 4; DB 5; Length 58;  
Best Local Similarity 100.0%, Fred. No. 2e+02;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5  
| | | |  
DB 11 VRMH 14

RESULT 13

US-10-057-498-11626

Sequence 11626, Application US/10007498

GENERAL INFORMATION:  
APPLICANT: Mitcham, Jennifer  
APPLICANT: Skeiky, Yasir  
APPLICANT: Persing, David  
APPLICANT: Bhutia, Ajay  
APPLICANT: Maisonneuve, Jean Francois  
APPLICANT: Zhang, Yanni  
APPLICANT: Jey, Chyian  
APPLICANT: Jey, William  
APPLICANT: Benson, Darin  
APPLICANT: Jones, Robert  
APPLICANT: Carter, Darick  
APPLICANT: Baruch, Brenda  
APPLICANT: Douglas, John  
TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Auto V  
FILE REFERENCE: 210121-514  
CURRENT APPLICATION NUMBER: US/10/057,498  
CURRENT FILING DATE: 2001-04-20  
NUMBER OF SEQ ID NOS: 23212  
SEQ ID NO 11626  
LENGTH: 58  
TYPE: PRT  
ORGANISM: Propionl acnes  
US 10-057-498-11626

Query Match 66.7%; Score 4; DB 5; Length 58;  
Best Local Similarity 100.0%, Fred. No. 2e+02;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5  
| | | |  
DB 11 VRMH 14

RESULT 14

PCT US02.32727-24772

Sequence 24772, Application PC/US0232727

GENERAL INFORMATION:  
APPLICANT: Mitcham, Jennifer  
APPLICANT: Skeiky, Yasir  
APPLICANT: Persing, David  
APPLICANT: Bhutia, Ajay  
APPLICANT: Maisonneuve, Jean Francois  
APPLICANT: Zhang, Yanni  
APPLICANT: Jey, Chyian  
APPLICANT: Jey, William  
APPLICANT: Benson, Michael  
APPLICANT: Benson, Darin  
APPLICANT: Jones, Robert  
APPLICANT: Carter, Darick  
APPLICANT: Baruch, Brenda  
APPLICANT: Douglas, John  
TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Auto V  
FILE REFERENCE: 210121-51401  
CURRENT APPLICATION NUMBER: PCT/US02/32727  
CURRENT FILING DATE: 2002-10-11  
NUMBER OF SEQ ID NOS: 39992  
SEQ ID NO 24772  
LENGTH: 64  
TYPE: PRT  
ORGANISM: Propionl acnes  
PCT US02.32727-24772

Query Match 66.7%; Score 4; DB 1; Length 64;  
Best Local Similarity 100.0%, Fred. No. 2e+02;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5  
| | | |  
DB 11 VRMH 14

RESULT 15

US-09-978-825-24772

Sequence 24772, Application US/09079825

GENERAL INFORMATION:

APPLICANT: Mitcham, Jennifer

APPLICANT: Skeiky, Yasir

APPLICANT: Persing, David

APPLICANT: Bhatia, Ajay

APPLICANT: Maisonneuve, Jean Francois

APPLICANT: Zhang, Yanni

APPLICANT: Wang, Siqing

APPLICANT: Jen, Shyian

APPLICANT: Lodes, Michael

APPLICANT: Benson, Darin

APPLICANT: Jones, Robert

APPLICANT: Carter, Darrick

APPLICANT: Barth, Brenda

APPLICANT: Douglass, John

TITLE OF INVENTION: Compositions and Methods for the Therapy and Diagnosis of Acne V

FILE REFERENCE: 210121.51401

CURRENT APPLICATION NUMBER: US/09/978,825

CURRENT FILING DATE: 2003-01-29

NUMBER OF SEQ ID NOS: 30942

SEQ ID NO: 24772

LENGTH: 64

TYPE: PRT

ORGANISM: Pseudomonas

US-09-978-825-24772

Query Match 66.7%; Score 4; DB 5; Length 64;

Best Local Similarity 100.0%; Pred. No. 2,28+02;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRMH 5

Db 2 VRMH 5

Search completed: February 20, 2003, 11:42:48  
Job time: 17.6667 secs







A:Cross-References: EMBL:Y90200, NID:340018, PIRN:PA016026.1, PIR:3134248  
 R/Borges, A.; Hawkins, C.F.; Packman, L.C.; Penham, P.N.  
 Eur. J. Biochem. 194, 95-102, 1990  
 A:Title: Cloning and sequence analysis of the genes encoding the dihydrofolate acetyl  
 Bacillus stearothermophilus.  
 A:Reference number: 146887  
 A:Accession: T46887  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-95 <BOR>  
 A:Cross-References: EMBL:Y90200, NID:340018, PIRN:PA016026.1, PIR:3134248  
 A:Experimental source: strain NC1503  
 C:Superfamily: Methanobacterium thermophilum conserved hypothetical protein MTH10

Query Match 66.7%; Score 4; DB 2; Length 95;  
 Best Local Similarity 100.0%; Pred.No.1:2e+02;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRWH 5  
 ||||  
 DB 5 VRWH 8

RESULT 8  
 AG3064  
 Hypothetical protein Acetivibrio (imported) - Acetivibrio tumefaciens (strain DSM, type strain)  
 C:Species: Acetivibrio tumefaciens  
 C:Date: 11-Jan-2002 #Sequence\_revision 11-Jan-2002 #text\_Change 11-Jan-2002  
 C:Accession: AG3064  
 R/Moore, J.W.; Gerbasi, R.A.; Kari, P.; Marks, D.; Chen, J.; Wood, G.F.; Chen, Y.; Wu, L.  
 et al.; G. Miller, W.; Grant, P.; Gonthier, P.; Kuyavin, T.; Levy, P.; Li, M.; McLeod,  
 J.; Karp, P.; Romero, P.; Zhang, S.  
 Science 294, 2317-2323, 2001  
 A:Authors: Yoon, H.; Tan, Y.; Riddle, P.; Tong, M.; Krespan, W.; Perry, M.; Gordon Farm,  
 ster, E.W.  
 A:Title: The Genome of the Natural Genetic Engineer Acetivibrio tumefaciens DSM  
 A:Reference number: AB2577; PMID:11743193  
 A:Accession: AG3064  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-98 <MIP>  
 A:Cross-References: EMBL:AF096093, PIRN:PA144933.1, PIR:31742997, GPCR:G000187  
 A:Experimental source: strain C58 (Dupont)  
 C:Genetics:  
 A:Gene: Atn4133  
 A:Map position: linear chromosome

Query Match 66.7%; Score 4; DB 2; Length 98;  
 Best Local Similarity 100.0%; Pred.No.1:2e+02;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VRWH 5  
 ||||  
 DB 3 VRWH 6

RESULT 9  
 A99800  
 Hypothetical protein Ecol369 (imported) - Escherichia coli (strain O157:H7, substrain PI  
 C:Species: Escherichia coli  
 C:Date: 10-Jul-2001 #Sequence\_revision 10-Jul-2001 #text\_Change 10-Jul-2001  
 C:Accession: A99800  
 R/Hayashi, T.; Makino, K.; Ohtsuki, M.; Furukawa, K.; Ishii, F.; Yokoyama, F.; Han, C.G.;  
 Gotohara, N.; Yasunaga, T.; Kohara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.  
 DNA Res. 8, 11-22, 2001  
 A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and gene  
 A:Reference number: A99800; MIMD:3115031; PMID:11258796  
 A:Accession: A99800  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-102 <HAY>  
 A:Cross-References: GB:BA000007; PIRN:PA934792.1, PIR:31368009, GSPDB:G00014  
 A:Experimental source: strain O157:H7, substrain RMD 0509952

C:Genetics:  
 A:Gene: ECS1369

Query Match 66.7%; Score 4; DB 2; Length 102;  
 Best Local Similarity 100.0%; Pred.No.1:3e+02;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 RMWF 6  
 ||||  
 DB 39 RMWF 42

RESULT 10  
 T27486  
 Hypothetical protein Y9C2UA.2, Caenorhabditis elegans  
 C:Species: Caenorhabditis elegans  
 C:Date: 10-Nov-1999 #Sequence\_revision 10-Nov-1999 #text\_Change 04-Mar-2000  
 C:Accession: T27486  
 R/Smye, R.  
 submitted to the EMBL Data Library, December 1998  
 A:Reference number: 220374  
 A:Accession: T27486  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-103 <WIL>  
 A:Cross-References: EMBL:AC009491, PIRN:PA101010.1, NID:340018, PIR:3134248  
 A:Experimental source: clone Y9C2UA  
 C:Genetics:  
 A:Gene: CESF-Y9C2UA.2  
 A:Map position 2  
 A:introns: 21/2  
 C:Superfamily: Caenorhabditis elegans hypothetical protein Y9C2UA.2

Query Match 66.7%; Score 4; DB 2; Length 103;  
 Best Local Similarity 100.0%; Pred.No.1:3e+02;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 RMWF 6  
 ||||  
 DB 53 RMWF 56

RESULT 11  
 T44499  
 Hypothetical protein (imported) Thermomonospora fusca (fragment)  
 C:Species: Thermomonospora fusca  
 C:Date: 21-Jan-2000 #Sequence\_revision 21-Jan-2000 #text\_Change 21-Jan-2000  
 C:Accession: T44499  
 R/Twin, D.L.; Chang, S.; Wilson, D.P.  
 submitted to the EMBL Data Library, April 1999  
 A:Description: Characterization of a Thermomonospora fusca family 42 serine protease  
 A:Reference number: 222783  
 A:Accession: T44499  
 A:Status: preliminary; translated from GB/EMBL/DBJ  
 A:Molecule type: DNA  
 A:Residues: 1-106 <IRW>  
 A:Cross-References: EMBL:AF144563; PIRN:AA09060.1  
 A:Experimental source: strain YX

Query Match 66.7%; Score 4; DB 2; Length 106;  
 Best Local Similarity 100.0%; Pred.No.1:3e+02;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRW 4  
 ||||  
 DB 14 WVRW 17

RESULT 12  
 MNXRI8  
 sigma IBSN protein - reovirus type 1 (strain Lang)  
 N:Alternative names: nonstructural protein  
 C:Species: reovirus type 1





GenCore version 5.1.3  
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OM protein - protein search, using sw model

Run on: February 20, 2003, 11:20:17, Search time: 11.133 seconds  
(without alignments)  
21,958 Million cell updates/sec

Title: US-09-912-414-2

Perfect score: 6

Sequence: 1 WVRMHF 6

Scoring table: OLIGODX  
Gapop 60.0, Gapext 60.0

Searched: 112892 seqs, 41476328 residues

Word size: 0

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summaries

Database: SwissProt\_40.0

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No	Score	Query Match length	DB ID	Description
1	83.3	373	GP45_HUMAN	Gq45(1) homo sapien
2	83.3	373	GP45_MOUSE	Gq45(1) mus musculu
3	83.3	794	AFSK_STRPO	P54741 streptomyc
4	83.3	807	AFSK_STRGR	P54742 streptomyc
5	66.7	95	VPDA_BACST	P21878 bacillus st
6	66.7	119	VSIS_PEOVL	P07938 reovirus (t
7	66.7	149	GLH1_MOPMP	P01197 moritatis mo
8	66.7	149	GLH1_PETMA	P08467 petromyzon
9	66.7	149	GLH2_MOPMP	P01198 moritatis mo
10	66.7	149	GLH2_PETMA	P01193 petromyzon
11	66.7	149	GLH3_MOPMP	P01196 moritatis mo
12	66.7	149	GLH3_PETMA	P08968 petromyzon
13	66.7	149	GLH5_PETMA	P08298 petromyzon
14	66.7	149	GIR_FAMFI	P08297 petromyzon
15	66.7	170	GRP4_YEAST	P31267 saccharomyc
16	66.7	195	TNF7_MOUSE	P05237 mus musculu
17	66.7	197	DSBE_XYLEA	O9pan4 xyella fas
18	66.7	208	YGED_BACSU	P54449 bacillus su
19	66.7	215	HPGP_POVIN	P49329 hae tauris
20	66.7	223	PGC2_HUMAN	C01173 homo sapien
21	66.7	244	YHGA_CABEL	P08067 caenorhathi
22	66.7	286	PURU_FOPSP	Q46339 corneabac
23	66.7	295	MOPA_PSPPI	Q02198 pseudomonas
24	66.7	298	YHIV_ECOLI	P33143 escherichia
25	66.7	310	DNIL_MYCGE	P47248 mycoplasma
26	66.7	323	ASTE_ECOLI	P36235 escherichia
27	66.7	325	HL76_HGOMA	P16712 human cytom
28	66.7	338	PDXA_PALSO	P58714 taliscoria s
29	66.7	403	YV67_MYCOT	Q56480 rhizobium s
30	66.7	403	YV4H_PHSIN	P55480 rhizobium s
31	66.7	418	PHAA_BACPH	P08219 bacillus ha
32	66.7	422	TERI_PSPPI	P53398 bacillus ha
33	66.7	436	GQ45_MOUSE	P08467 mus musculu

34	4	66.7	440	1	SOC4_HUMAN	G0455 homo sapien
35	4	66.7	450	1	YVE3_PSEAB	G0453 pseudomonas
36	4	66.7	462	1	TRIM_YPACT	P07316 escherichia
37	4	66.7	472	1	PRB_ESCCE	P21948 escherichia
38	4	66.7	475	1	ETS6_PPOWE	P29726 drosophila
39	4	66.7	486	1	YDF1_ECOLI	P12260 escherichia
40	4	66.7	507	1	TRAG_PSEAE	G07541 pseudomonas
41	4	66.7	508	1	TRAT_CHEHE	G16993 chelidonia
42	4	66.7	511	1	FAST_MOUSE	G01189 mus musculu
43	4	66.7	543	1	FAST_HUMAN	G14256 homo sapien
44	4	66.7	583	1	STS_HUMAN	P08842 homo sapien
45	4	66.7	611	1	SNF1_CANDA	G00372 candida gla

## ALIGNMENTS

RESULT 1	GP45_HUMAN	STANDARD:	PRT:	372 AA.
AC	Q95Y33:			
DT	15-JUN-2002 (rel. 41, last sequence update)			
DT	15-JUN-2002 (rel. 41, last annotation update)			
DE	Probable G-protein-coupled receptor GP45 (PSP24-alpha) (PSP24-1).			
GN	GP45.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eumetazoa; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.			
OX	NCBI_TaxID=9606;			
RN	(1)			
RP	SEQUENCE FROM N.A.			
FY	MEDLERE:9115652; PubMed 1027181;			
EA	Marichev A, Sawadsky M, Rajewsky T, Cheng F, Heng H-H, Brock T, Im D-S, Lynch K.P., George S.P., O'Dowd B.F.;			
PA	"Discovery of three novel orphan G-protein-coupled receptors.";			
RL	Genomics 56:12-21(1999).			
CC	- FUNCTION: Orphan receptor. May play a role in brain function.			
CC	- SUBCELLULAR LOCATION: Integral membrane protein.			
CC	- TISSUE SPECIFICITY: Expressed in brain; detected in the basal forebrain, frontal cortex, and caudate, but not in thalamus, hippocampus, or putamen.			
CC	- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation.			
CC	the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see <a href="http://www.ebi.ac.uk/submit/">http://www.ebi.ac.uk/submit/</a> or send an email to <a href="mailto:license@ebi.ac.uk">license@ebi.ac.uk</a> ).			
DR	EMBL: A019066; A019066.1; -			
DR	GenBank: AF091453; SPP45.			
DR	MIM: 604838; -			
DR	InterPro: IPR000276; GPCR_Fhdpsn.			
DR	Pfam: PF00001; 7tm1.1.			
DR	PRINTS: P000237; GPCRPhosphn.			
DR	PROSITE: P000237; G-PROTEIN RECEPTOR_F1; FALSE_NEG.			
EF	PROSITE: PS00062; G-PROTEIN RECEPTOR_F2; 1.			
FM	G-protein coupled receptor; Transmembrane protein; G-protein.			
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FT	TRANSMEM 39 59			
FT	DOMAIN 60 75			
FT	TRANSMEM 76 96			
FT	DOMAIN 97 109			
FT	TRANSMEM 110 136			
FT	DOMAIN 137 149			
FT	TRANSMEM 150 172			
FT	DOMAIN 173 198			
FT	TRANSMEM 199 219			
FT	DOMAIN 220 265			
FT	TRANSMEM 269 289			

[illegible]

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Query Match:      83.38;  Score 5;  DB 1;  Length 372;
Best Local Similarity 100.0%;  Pred No 4.4;
Matches 5;  Conservative 0;  Mismatches 0;  Indels 0;  Gaps 0

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QY	2	VERB	6
	1	VERB	6
	1	VERB	6
	1	VERB	6
QB	98	VERB	102

[illegible][illegible]

The European Information Institute for Security and Humanitarian Affairs (EIISS) is a non-profit organization established in 1992. It is a member of the European Union and the United Nations. The EIISS is a leading international organization in the field of security and humanitarian affairs. It provides a wide range of services, including research, training, and information. The EIISS is also a member of the European Union's Security and Humanitarian Affairs Committee.

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76	DOWN	246	214	EXPANDED (POTENTIAL)
77	DOWN	247	215	EXPANDED (POTENTIAL)
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98 VPMHF 102

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D1 15-001-001 (ref. 49) Last sequence update)  
 D2 15-001-002 (ref. 47) Last annotation update)  
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CC SUBCLONING FROM N.A.  
 CT STRAIN A3(2)?  
 CX MEL:10E 3424108, Feb93 807354,  
 XX Matsumoto A., Hong S.K., Shikata H., Hattinouchi S., Beppu T.;  
 RT "Phosphorylation of the ATR Kinase Involved in Secondary and  
 RT 1. Antiproliferative Effects by a Catalytic Type Protein Kinase.";  
 RL Gene 146:147-56(1994).  
 RN 121  
 RN SEQUENCE FROM N.A.  
 RP STRAIN A3(2)?  
 RC MEL:10E 3424108, Feb93 807354/  
 RX Hattinouchi S., Beppu T., Horinouchi S.;  
 RA "Phosphorylation of the ATR Kinase Involved in Secondary and  
 RA 1. Antiproliferative Effects by a Catalytic Type Protein Kinase  
 RI of 5' (nucleotide 1570)."  
 RI Gene 146:147-56(1994)

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CC EMBL: D45382; GAA09209.1;  
 DR FPMU; A1392174; CAC08432.1;  
 DR FPMU; A1392174; CAC08432.1;  
 DR Phosphite; P54741;  
 DR InterPro: IPR000719; Euk kinase.  
 DR InterPro: IPR002280; Serine/threonine  
 DR InterPro: IPR001245; Tyk kinase.  
 DR Pfam: PF00069; Pkinase; 1.  
 DR ProDom: PD00001; Euk Pkinase; 1.  
 DR SMART: SM00222; S\_TKc; 1.  
 DR SMART: SM00219; TykKc; 1.  
 DR PROSITE: PS00107; PROTEIN\_KINASE\_ATP\_1  
 DR PROSITE: PS00109; PROTEIN\_KINASE\_ATP\_1  
 DR PROSITE: PS00111; PROTEIN\_KINASE\_DOM; 1.  
 KW Transferrase; Serine/threonine-protein kinase; ATP binding;  
 KW Phosphorylation; Complete from dom.  
 FT DOMAIN 16 271 PROTEIN KINASE.  
 FT NR BIND 22 30 ATP (BY SIMILARITY).  
 FT BINDING 44 44 ATP (BY SIMILARITY).  
 FT ACT SITE 138 138 BY SIMILARITY.  
 SQ SEQUENCE 744 AA; 84747 MW; 4FE5E415E5F5E C0664;

Query Match 63.3%; Score 5; DB 1; Length 709;  
 Best Local Similarity 100.0%; Pred. No. 9.6;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 VRWHF 6  
 Db 628 VRWHF 632

RESULT 4  
 AFSK\_STRCP STANDARD; PRT; 807 AA.  
 ID AFSK\_STRCP  
 AC P4742;  
 DT 01-OCT-1996 (rel. 34, Created)  
 DT 01-OCT-1996 (rel. 34, Last sequence update)  
 DT 30-MAY-2000 (rel. 39, Last annotation update)  
 GN Serine/threonine protein kinase atsk (E02771 -)  
 GN AFSK.  
 CC Streptococcus glaucus.  
 CC Bacteria; Actinobacteria; Actinobacteriales; Actinobacteriia;  
 CC Actinomycetales; Streptococcaceae; Streptococcaceae; Streptococcus  
 CC NCBI\_TaxID:911;  
 RN NCBI\_TaxID:911;  
 PP SEQUENCE FROM N A  
 PY MFEU; 0618000, PubMed:943767.  
 PA Ueda K., Uneyama T., Beppu T., Horinouchi S.,  
 RT The aerial mycelium defective phenotype of *Streptococcus glaucus*  
 RT resulting from A-factor deficiency is suppressed by a Ser/thr kinase  
 RT of *S. coelicolor* A3(2).  
 RL Gene 169;91-95(1996).  
 CC -1- FUNCTION: INVOLVED IN THE REGULATION OF SECONDARY METABOLISM BY  
 CC PHOSPHORYLATION, ON BOTH SER AND THR. THE AFSK GLOBAL REGULATORY  
 CC PROTEIN INVOLVED IN THE CONTROL OF SECONDARY METABOLISM (BY  
 CC SIMILARITY).  
 CC -1- PTM: AUTOPHOSPHORYLATED AT THR AND SER RESIDUES (BY SIMILARITY).  
 CC -1- SIMILARITY: PELOWS TO THE SER/THR FAMILY OF PROTEIN KINASES  
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CC EMBL: D45246; BAA08203.1;  
 DR Phosphite; P54742;

DR InterPro: IPR000719; Euk kinase.  
 DR InterPro: IPR002280; Ser/thr\_kinase.  
 DR Pfam: PF00069; Pkinase; 1.  
 DR ProDom: PD00001; Euk Pkinase; 1.  
 DR PROSITE: PS00107; PROTEIN\_KINASE\_ATP\_1  
 DR PROSITE: PS00109; PROTEIN\_KINASE\_ATP\_1  
 DR PROSITE: PS00111; PROTEIN\_KINASE\_DOM; 1.  
 KW Transferrase; Serine/threonine-protein kinase; ATP-binding;  
 KW Phosphorylation.  
 KM Phosphorylation.  
 FT DOMAIN 16 272 PROTEIN KINASE.  
 FT NR BIND 22 30 ATP (BY SIMILARITY).  
 FT BINDING 44 44 ATP (BY SIMILARITY).  
 FT ACT SITE 138 138 BY SIMILARITY.  
 SQ SEQUENCE 807 AA; 84747 MW; 4FE5E415E5F5E C0664;

Query Match 63.3%; Score 5; DB 1; Length 807;  
 Best Local Similarity 100.0%; Pred. No. 8.7;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 VRWHF 6  
 Db 635 VRWHF 639

RESULT 5  
 YPDA\_BACST STANDARD; PRT; 95 AA.  
 ID YPDA\_BACST  
 AC F21879;  
 DT 01-MAY-1991 (rel. 18, Created)  
 DT 01-MAY-1991 (rel. 18, Last sequence update)  
 DT 16-OCT-2001 (rel. 40, Last annotation update)  
 DE Hypothetical protein in pDNA 5' region (ORF1) (Fragment).  
 DE Bacillus stearothermophilus.  
 CC Bacteria; Firmicutes; Bacillales; Geobacillus.  
 CC NCBI\_TaxID:1422;  
 RN NCBI\_TaxID:1422;  
 PP SEQUENCE FROM N A  
 RC STRAIN:YCA 1503,  
 RX MEDLINE:50345939, PubMed:2206674,  
 RA Hawkins C.F., Borges A., Perham R.N.;  
 PT cloning and sequence analysis of the genes encoding the alpha and  
 PT beta subunits of the E1 component of the pyruvate decarboxylase  
 PT multienzyme complex of *Bacillus stearothermophilus*.  
 RL Eur. J. Biochem. 191:337-346(1990).  
 CC -1- SIMILARITY: PELOWS TO THE SER/THR FAMILY OF PROTEIN KINASES  
 CC (B.SUBTILIS) FAMILY.  
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CC EMBL: X53560; CAA37626.1;  
 DR PIP; S10796; S10796.  
 DR InterPro: IPR001454; Hydrolase; Hydrolase.  
 DR InterPro: IPR000150; Hypothet\_cof.  
 DR Pfam: PF00702; Hydroxylase; 1.  
 DR PROSITE: PS01229; COF\_1; EAF1AL.  
 DR PROSITE: PS01229; COF\_2; 1.  
 KW Hypothetical protein.  
 FT NON TYP 1  
 SQ SEQUENCE 95 AA; 10441 MW; 811E9487321A74E3 C0664;

Query Match 66.7%; Score 4; DB 1; Length 95;  
 Best Local Similarity 100.0%; Pred. No. 27;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 VRWH 5  
 Db 5 VRWH 8



DT 15-JUN-2002 (rel 4), last annotation update)  
 DE Globin II.  
 OS Moradacia mordax (Southern hemisphere lamprey)  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Hyperoartia;  
 OC Petromyzontiformes; Petromyzontidae; Moradacia.  
 OX NCBI\_TaxID=7755;  
 RN [1]  
 RP SEQUENCE.  
 RX MEDLINE:3124417, PubMed:303605;  
 RA Hombrades I., Vidal Y., Podewald K., Braunitzer G., Neužil F.,  
 RT "The primary structure of the hemoglobins of a southern hemisphere  
 lamprey (Moradacia mordax, Cyclostomata)".  
 RL Biol. Chem. Hoppe-Seyler 372:49-56(1991).  
 CC -1- SUBUNIT: Monomer.  
 CC -1- SIMILARITY: BELONGS TO THE GLOBIN FAMILY.  
 DR PIR: S13459; S13459.  
 DR HSSP: P02208; 2LHB.  
 DR InterPro: IPR000971; Globin.  
 DR Pfam: PF00042; Globin\_1.  
 DR PROSITE: PS01033; GLOBIN; 1.  
 KM Heme: Oxygen transport; Transport.  
 SQ SEQUENCE 149 AA; 14908 MW; 1416044566 CP064;  
 Query Match 66.7%; Score 4; DB 1; Length 149;  
 Best Local Similarity 100.0%; Pred. No. 40;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 2 VRWH 5  
 Db 70 VRWH 73  
 RESULT 10  
 GLB3\_PETMA STANDARD; PRT; 149 AA.  
 ID GLB3\_PETMA  
 AC 091913;  
 DT 16-OCT-2001 (rel 40, Created)  
 DT 16-OCT-2001 (rel 40, Last sequence update)  
 DE Globin II (Hemoglobin PM1).  
 OS Petromyzon marinus (Sea lamprey).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Hyperoartia;  
 OC Petromyzontiformes; Petromyzontidae; Petromyzon.  
 OX NCBI\_TaxID=7757;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE:20350908; PubMed:10788466;  
 RA Qiu Y., Maillett D.H., Knapp J., Olson J.S., Piggs A.F.;  
 RT "Lamprey hemoglobin: Structural basis of the Bohr effect".  
 RL J. Biol. Chem. 275:13517-13528(2000).  
 CC -1- SIMILARITY: BELONGS TO THE GLOBIN FAMILY.  
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 CC  
 DR EMBL: AF248645; AAB67186.1;  
 DR HSSP: P02208; 2LHB.  
 DR InterPro: IPR000971; Globin.  
 DR Pfam: PF00042; Globin\_1.  
 DR PRINTS: PR00613; MYOGLOBIN.  
 DR PROSITE: PS01033; GLOBIN; 1.  
 KM Heme: Oxygen transport; Transport.  
 FT INIT MET 0  
 SQ SEQUENCE 149 AA; 14911 MW; 17830643241746 CP064;  
 Query Match 66.7%; Score 4; DB 1; Length 149;  
 Best Local Similarity 100.0%; Pred. No. 40;

Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 2 VRWH 5  
 Db 70 VRWH 73  
 RESULT 11  
 GLB3\_MORDE STANDARD; PRT; 149 AA.  
 ID GLB3\_MORDE  
 AC P21199;  
 DT 01-MAY-1991 (rel 18, Created)  
 DT 01-MAY-1991 (rel 18, Last sequence update)  
 DT 15-JUN-2002 (rel 41, Last annotation update)  
 DE Globin III.  
 OS Moradacia mordax (Southern hemisphere lamprey).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Hyperoartia;  
 OC Petromyzontiformes; Petromyzontidae; Moradacia.  
 OX NCBI\_TaxID=7755;  
 RN [1]  
 RP SEQUENCE.  
 RX MEDLINE:3124417, PubMed:303605;  
 RA Hombrades I., Vidal Y., Podewald K., Braunitzer G., Neužil F.,  
 RT "The primary structure of the hemoglobins of a southern hemisphere  
 lamprey (Moradacia mordax, Cyclostomata)".  
 RL Biol. Chem. Hoppe-Seyler 372:49-56(1991).  
 CC -1- SUBUNIT: Monomer.  
 CC -1- SIMILARITY: BELONGS TO THE GLOBIN FAMILY.  
 DR PIR: S13460; S13460.  
 DR HSSP: P02208; 2LHB.  
 DR InterPro: IPR000971; Globin.  
 DR Pfam: PF00042; Globin\_1.  
 DR PROSITE: PS01033; GLOBIN; 1.  
 KM Heme: Oxygen transport; Transport.  
 SQ SEQUENCE 149 AA; 14908 MW; 21603504456666 CP064;  
 Query Match 66.7%; Score 4; DB 1; Length 149;  
 Best Local Similarity 100.0%; Pred. No. 40;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 2 VRWH 5  
 Db 70 VRWH 73  
 RESULT 12  
 GLB3\_PETMA STANDARD; PRT; 149 AA.  
 ID GLB3\_PETMA  
 AC P09568;  
 DT 01-MAY-1989 (rel 10, Created)  
 DT 01-MAY-1989 (rel 10, Last sequence update)  
 DT 16-OCT-2001 (rel 40, Last annotation update)  
 DE Globin III.  
 OS Petromyzon marinus (Sea lamprey).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Hyperoartia;  
 OC Petromyzontiformes; Petromyzontidae; Petromyzon.  
 OX NCBI\_TaxID=7757;  
 RN [1]  
 RP SEQUENCE.  
 RX MEDLINE:87184913, PubMed:3566915;  
 RA Hombrades I., Podewald K., Allard M., Neužil F., Braunitzer G.,  
 RT "Primary structure of the minor hemoglobins from the sea lamprey  
 (Petromyzon marinus, Cyclostomata)".  
 RL Biol. Chem. Hoppe-Seyler 369:145-154(1987).  
 CC -1- SUBUNIT: Monomer.  
 CC -1- MISCELLANEOUS: THIS IS ONE OF THE MINOR GLOBIN COMPONENT OF SEA  
 LAMPREY.  
 CC -1- SIMILARITY: BELONGS TO THE GLOBIN FAMILY.  
 DR PIR: P26042; B26042.  
 DR HSSP: P02208; 2LHB.  
 DR InterPro: IPR000971; Globin.  
 DR Pfam: PF00042; Globin\_1.  
 DR PROSITE: PS01033; GLOBIN; 1.



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ID   CBP4 YEAST          STANDARD;          PRT;          170 AA.
AC   P17267;
DT   01-OCT-1994 (Rel. 30, Created)
DT   01-OCT-1996 (Rel. 34, Last sequence update)
DT   16-OCT-2001 (Rel. 40, Last annotation update)
DE   CBP4 Protein, mitochondrial precursor.
GN   CBP4 OR YGR174C.
OS   Saccharomyces cerevisiae (Baker's Yeast).
OC   Eukaryota, Fungi, Ascomycota, Saccharomycotina, Saccharomycetes,
OC   Saccharmycetales, Saccharomycetaceae, Saccharomyces.
OX   NCBI_TaxID=4932;
RN   [1]
RP   SEQUENCE FROM N.A.
RX   MEDLINE=94142101; PubMed=8063753;
RA   Crivellone M.D.;
RT   "Characterization of CBP4, a new gene essential for the expression of
RT   ubiquinol-cytochrome c reductase in Saccharomyces cerevisiae.";
PL   J Biol Chem. 269(21):2084-2102(1994).
RN   [2]
RP   SEQUENCE FROM N.A.
RA   Hebling U., Hofmann B., Delius H.;
RA   Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
CC   -1- FUNCTION: ESSENTIAL FOR THE ASSEMBLY AND/OR STABILITY OF
CC   TRICLINOL-CYTOCHROME C REDUCTASE. IT HAS A DIRECT EFFECT ON THE
CC   CATALYTIC ACTIVITY OF THE P185F PROTEIN, COPE 4, COPE 5 AND
CC   APOCYTOCHROME P. IT MAY EITHER BE INVOLVED IN POST-TRANSLATIONAL
CC   MODIFICATION OF THE SUBUNITS OR IN THE ASSEMBLY OF THE ENZYME.
CC   -2- SUBCELLULAR LOCATION: MITOCHONDRIAL ASSOCIATED WITH THE INNER
CC   MEMBRANE.
CC   -----
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DR   EMBL; U10700; AAA61566.1; -
DR   EMBL; Z72950; CAA07220.1; -
DR   PIR; A53928; A53928.
DR   SCS; S0003406; CBP4.
FM   Mitochondrion; Transit peptide.
FT   TRANSIT 1
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Query Match At 1% Score 5; DB 15; Length 156;  
 Best Local Similarity 100.0%; Pred. No. 47;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 VWRMH 6  
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 DB 27 VWRMH 31

## RESULT 5

Q9M952 PRELIMINARY; PRT; 370 AA.

AC Q9M952; 01-OCT-2000 (TRENBLER) 15, Created;  
 DT 01-OCT-2000 (TRENBLER) 15, Last sequence update;  
 DT 01-JUN-2002 (TRENBLER) 21, Last annotation update;  
 DE F14023.22 protein;  
 GN F14023.22;  
 OS Arabidopsis thaliana (Mouse-ear cress);  
 OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;  
 OC Spermatophyta; Magnoliopsida; Eudicotyledons; Core eudicotyledons; Rosidae;  
 OC Euphorbiales; Brassicales; Brassicaceae; Arabidopsis  
 NC NCB1\_TaxID:3702;  
 RX (1)  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-CV, COLUMBIA;  
 RA Liu S., Yu G., Lee S., Sakano H., Bhavani A., Lee C., Toriumi M.,  
 RA Liu C., Chien J., Choi E., Gonzalez A., Huang P., Koo T., Li J.,  
 RA Liu A., Pham P., Vaynsberg M., Altschul H., Ruchler E., Chao Q., Conn L.,  
 RA Nguyen M., Palm C., Shinn P., Tambunga C., Davis P., Ecker J.,  
 RA Rostkowski J., Twell J., Shinn P.,  
 RA The sequence of BAC F14023 from Arabidopsis thaliana chromosome 1;  
 RL Submitted (NOV 1999) to the EMBL/GenBank/CCDB databases  
 RN (2)  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-CV, COLUMBIA;  
 RA Theologis A;  
 RL Submitted (MAR 2000) to the EMBL/GenBank/CCDB databases  
 RX (1) Similarity: MAP 26001 to the EMBL/GenBank/CCDB databases  
 DR EMBL: A0012654; AAF43237.1;  
 DR Interpro: IPR001480; WD40  
 DR Pfam: PF00440; WD40; 7  
 DR PRINTS: PR00129; GPROTEINRPT  
 DR SMART: SM00320; WD40; 7  
 DR PROSITE: PS00678; WD\_REPEATS\_1; UNKNOWN;  
 DR PROSITE: PS00682; WD\_REPEATS\_2; 7  
 DR PROSITE: PS00704; WD\_REPEATS\_REGION; 1  
 KW Repeat; WD repeat  
 SQ SEQUENCE 370 AA; 14774 MW; 716601F5403E15 CPO64;

Query Match At 1% Score 5; DB 19; Length 170;  
 Best Local Similarity 100.0%; Pred. No. 48;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 VWRMH 5  
 |||||  
 DB 151 VWRMH 155

## RESULT 6

Q43898 PRELIMINARY; PRT; 376 AA.

AC Q43898; 01-JUN-1998 (TRENBLER) 06, Created;  
 DT 01-JUN-1998 (TRENBLER) 06, Last sequence update;  
 DT 01-JUN-2002 (TRENBLER) 21, Last annotation update;  
 DE High-affinity lysophosphatidic acid receptor homolog;  
 OS Homo sapiens (Human);  
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Hominidae;  
 NC NCB1\_TaxID:9606;

RN (1)  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-BRAIN;  
 RA An S.;

RL Submitted (MAR-1997) to the EMBL/GenBank/CCDB databases  
 RF EMBL: U92642; AAB93844.1;  
 DR Interpro: IPR000276; GPCR\_Phodpsn.  
 DR Pfam: PF00001; 7tm\_1; 1.  
 DR PRINTS: PR00237; GPCRPHODPSN  
 DR PROSITE: PS00621; G\_PROTEIN\_RECEP\_F1\_1;  
 KW Receptor.  
 SQ SEQUENCE 376 AA; 4456 MW; 44C913B82C8430DF CPO64;

Query Match At 1% Score 5; DB 4; Length 376;  
 Best Local Similarity 100.0%; Pred. No. 48;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 VWRMH 6  
 |||||  
 DB 98 VWRMH 102

## RESULT 7

Q8RX04 PRELIMINARY; PRT; 407 AA.

AC Q8RX04; 01-JUN-2002 (TRENBLER) 21, Created;  
 DT 01-JUN-2002 (TRENBLER) 21, Last sequence update;  
 DT 01-JUN-2002 (TRENBLER) 21, Last annotation update;  
 DE Hypothetical 43 kDa protein.  
 GN A1671840.  
 OS Arabidopsis thaliana (Mouse-ear cress);  
 OC Eukaryota; Viridiplantae; Streptophyta; Eudicotyledons; Tracheophyta;  
 OC Spermatophyta; Magnoliopsida; Eudicotyledons; Core eudicotyledons; Rosidae;  
 OC Euphorbiales; Brassicales; Brassicaceae; Arabidopsis  
 NC NCB1\_TaxID:3702;  
 RX (1)  
 RP SEQUENCE FROM N.A.  
 RA Yamada K., Liu S.X., Sakano H., Pham P.K., Panth J., Chung M.K.,  
 RA Goldsmith A.F., Lee S.M., Quach H.L., Toriumi M., Yu G., Bowser L.,  
 RA Cantarel F., Chen H., Cheek P., Hayashikawa Y., Ishida T., Jones T.,  
 RA Kamuya A., Karlin-Neumann G., Kawai Y., Kim C., Lam P., Lin T.,  
 RA Miranda M., Narusaka M., Nguyen M., Palm C.J., Sakurai T., Satou M.,  
 RA Seki M., Shinn P., Southwick A., Shinozaki K., Shinozaki P., Davis P.W., Ecker J.P.,  
 RA Theologis A.;

PT "Arabidopsis full length cDNA clones."  
 RL Submitted (FEB 2002) to the EMBL/GenBank/CCDB databases.  
 DR EMBL: AY080732; AAL86002.1;  
 KW Hypothetical protein.  
 SQ SEQUENCE 407 AA; 43791 MW; 712D70ADD1D94833 CPO64;

Query Match At 1% Score 5; DB 10; Length 407;  
 Best Local Similarity 100.0%; Pred. No. 51;  
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 VWRMH 5  
 |||||  
 DB 160 VWRMH 164

## RESULT 8

Q97774 PRELIMINARY; PRT; 52 AA.

AC Q97774; 01-OCT-2001 (TRENBLER) 19, Created;  
 DT 01-OCT-2001 (TRENBLER) 19, Last sequence update;  
 DT 01-DEC-2001 (TRENBLER) 19, Last annotation update;  
 DE Hypothetical protein SP0077.  
 GN SP0077.  
 OS Streptococcus pneumoniae.  
 OC Bacteria; Firmicutes; Bacillus/Clostridium group; Bacillales;  
 OC Streptococcales; Streptococcus.  
 NC NCB1\_TaxID:1313;



KW Hypothetical protein.  
 FT NON TER  
 SO SEQUENCE 83 AA: 9486 MW: 9960798179441ED CRC64;

Query Match 66.7%; Score 4; DB 11; Length 83;  
 Best Local Similarity 100.0%; Pred No 2 50.0%;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 VRW 4  
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 Db 54 VRW 57

RESULT 13  
 Q96812 PRELIMINARY; PRT; 91 AA.

AC Q96812; PRELIMINARY; PRT; 91 AA.  
 DT 01-FEB-1997 (TREMBLrel 02, Created)  
 DT 01-FEB-1997 (TREMBLrel 02, Last sequence update)  
 DT 01-DEC-2001 (TREMBLrel 19, Last annotation update)  
 DE RNA binding protein.  
 OS Grapevine virus D.  
 OC Viruses; ssRNA positive strand viruses; ss- RNA stage, Vi. virus  
 ON NCPI\_TaxID=51617;  
 RN 11  
 RP SEQUENCE FROM N.A.  
 RA Abou Ghannem N., Salazar P., Minafra A., Buzkan N., Castellano M.A.,  
 RA Martelli G.P.;  
 RT "Properties of grapevine virus D, a novel putative "trichovirus";  
 RL J. Plant Pathol. 94:15-24(1997)  
 DR EMBL; Y07764; CAA69071.1; -  
 SO SEQUENCE 91 AA: 10456 MW: 7358255530448F CRC64;

Query Match 66.7%; Score 4; DB 12; Length 91;  
 Best Local Similarity 100.0%; Pred No. 2 70.0%;  
 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 VRW 4  
 ||||  
 Db 70 VRW 73

RESULT 14  
 Q8U0Y1 PRELIMINARY; PRT; 94 AA.

AC Q8U0Y1; PRELIMINARY; PRT; 94 AA.  
 DT 01-JUN-2002 (TREMBLrel 21, Created)  
 DT 01-JUN-2002 (TREMBLrel 21, Last sequence update)  
 DT 01-JUN-2002 (TREMBLrel 21, Last annotation update)  
 DE Hypothetical protein PF1450.  
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 FA Chumley F., Tingey S.V., Tomb J.-F., Gordon M.P., Olson M.V.,  
 FA Neeter E.W.;  
 FT "The genome of the natural genetic engineer Agrobacterium tumefaciens  
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KW Protein identification; signal transduction pathway; metabolic pathway;
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KW linoleic-acid-desaturase; oleic-desaturase; transgenic plant;

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OS Not specified.

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PR 08-APR-1999; 99US-0128714.

PR 16-APR-1999; 99US-0129845.

PR 19-APR-1999; 99US-0130077.

PR 21-APR-1999; 99US-0130449.

PR 23-APR-1999; 99US-0130510.

PR 23-APR-1999; 99US-0130891.

PR 28-APR-1999; 99US-0131440.

PR 30-APR-1999; 99US-0132048.

PR 04-MAY-1999; 99US-0132484.

PR 05-MAY-1999; 99US-0132485.

PR 06-MAY-1999; 99US-0132486.

PR 07-MAY-1999; 99US-0132487.

PR 11-MAY-1999; 99US-0132863.

PR 14-MAY-1999; 99US-0134218.

PR 14-MAY-1999; 99US-0134219.

PR 14-MAY-1999; 99US-0134221.

PR 18-MAY-1999; 99US-0134370.

PR 19-MAY-1999; 99US-0134768.

PR 20-MAY-1999; 99US-0134941.

PR 21-MAY-1999; 99US-0135124.

PR 24-MAY-1999; 99US-0135438.

PR 25-MAY-1999; 99US-0136021.

PR 27-MAY-1999; 99US-0136192.

PR 28-MAY-1999; 99US-0136782.

PR 01-JUN-1999; 99US-0137222.

PR 03-JUN-1999; 99US-0137528.

PR 04-JUN-1999; 99US-0137602.

PR 07-JUN 1999; 99US-0137724.

PR 08-JUN-1999; 99US-0138094.

PR 10-JUN-1999; 99US-0138540.

PR 10-JUN-1999; 99US-0138847.

PR 14-JUN-1999; 99US-0139119.

PR 16-JUN-1999; 99US-0139452.

PR 17-JUN-1999; 99US-0139453.

PR 17-JUN-1999; 99US-0139482.

PR 18-JUN-1999; 99US-0139454.

PR 18-JUN-1999; 99US-0139455.

PR 18-JUN-1999; 99US-0139456.

PR 18-JUN-1999; 99US-0139457.

PR 18-JUN-1999; 99US-0139458.

PR 18-JUN-1999; 99US-0139459.

PR 18-JUN-1999; 99US-0139460.

PR 18-JUN-1999; 99US-0139461.

PR 18-JUN-1999; 99US-0139462.

PR 18-JUN-1999; 99US-0139463.

PR 18-JUN-1999; 99US-0139750.

PR 18-JUN-1999; 99US-0139761.

PR 21-JUN-1999; 99US-0139817.

PR 22-JUN-1999; 99US-0139849.

PR 23-JUN-1999; 99US-0140351.

PR 24-JUN-1999; 99US-0140354.

PR 24-JUN-1999; 99US-0140695.

Query Match aa 98; Score 40; PP 15; length 419;

Best Local Similarity 81%; Prod No 81;

Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 WVRWHP 6

DB 257 WVRWHP 260

# RESULT 14

AA060508 standard; Protein; 448 AA.

AC AAG05808;

DT 17-OCT-2000 (first entry)

DE Arabidopsis thaliana protein fragment SEQ ID NO: 3152

KW Protein identification; signal transduction pathway; metabolic pathway;

XX hybridization assay; genetic mapping; gene expression control; promoter;







PR 04-SEP-1999; 99US-0155659;  
PR 28-SEP-1999; 99US-0155659;  
PR 29-SEP-1999; 99US-0155659;  
PR 04-OCT-1999; 99US-0157117;  
PR 04-OCT-1999; 99US-0157117;  
PR 04-OCT-1999; 99US-0157117;  
PR 04-OCT-1999; 99US-0157865;  
PR 07-OCT-1999; 99US-0158029;  
PR 08-OCT-1999; 99US-0158029;  
PR 12-OCT-1999; 99US-0158029;  
PR 13-OCT-1999; 99US-0158293;  
PR 13-OCT-1999; 99US-0158293;  
PR 13-OCT-1999; 99US-0158294;  
PR 13-OCT-1999; 99US-0158295;  
PR 14-OCT-1999; 99US-0158329;  
PR 14-OCT-1999; 99US-0158329;  
PR 14-OCT-1999; 99US-0158330;  
PR 14-OCT-1999; 99US-0158331;  
PR 14-OCT-1999; 99US-0158337;  
PR 14-OCT-1999; 99US-0158338;  
PR 14-OCT-1999; 99US-0158338;  
PR 21-OCT-1999; 99US-0160741;  
PR 21-OCT-1999; 99US-0160741;  
PR 21-OCT-1999; 99US-0160767;  
PR 21-OCT-1999; 99US-0160768;  
PR 21-OCT-1999; 99US-0160770;  
PR 21-OCT-1999; 99US-0160814;  
PR 21-OCT-1999; 99US-0160815;  
PR 22-OCT-1999; 99US-0160980;  
PR 22-OCT-1999; 99US-0160981;  
PR 22-OCT-1999; 99US-0160989;  
PR 25-OCT-1999; 99US-0161404;  
PR 25-OCT-1999; 99US-0161405;  
PR 25-OCT-1999; 99US-0161406;  
PR 26-OCT-1999; 99US-0161359;  
PR 26-OCT-1999; 99US-0161360;  
PR 26-OCT-1999; 99US-0161361;  
PR 28-OCT-1999; 99US-0161920;  
PR 28-OCT-1999; 99US-0161992;  
PR 28-OCT-1999; 99US-0161993;  
PR 29-OCT-1999; 99US-0162142;

Query Match 88.9%; Score 40; DH 21; Length 448;  
Percent Similarity 83.1%;  
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 WVRWF 6  
DB 257 WVRWF 262

Search Completed: February 20, 2003, 10:47:34  
Total time: 38.3313 secs

GenScore version 5.1.1  
Copyright (c) 1991 - 2003 Copyright Ltd.

OM protein - protein search, using sw model

Run on: February 20, 2003, 10:46:13, Search time 12 seconds

(without alignments)  
48,067 Million cell updates/sec

Title: US-09-912-414-2

Perfect score: 45

Sequence: 1 WPMWP 6

Scoring table: BLOSUM62, Gapop 10.0, Gapext 0.5

Searched: 293224 seqs, 9613442 residues

Total number of hits satisfying chosen parameters: 293224

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: PIP 71.1  
1: p1r1:  
2: p1r2:  
3: p1r3:  
4: p1r4:

Pred. No. is the number of results potentially changed by a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	40	88.9	448	2	T08136
2	40	88.9	448	2	probable omega-6
3	39	86.7	293	2	hypothetical prote
4	39	86.7	370	2	hypothetical prote
5	39	86.7	482	2	hypothetical prote
6	38	84.4	477	1	catalase (EC 1.11.
7	38	84.4	822	1	brain-derived neur
8	37	82.2	591	2	probable protein 5
9	36	80.0	185	2	hypothetical prote
10	36	80.0	230	2	hypothetical prote
11	36	80.0	262	2	hypothetical prote
12	36	80.0	266	2	probable peroxide
13	36	80.0	266	2	probable aliphatic
14	36	80.0	394	2	hypothetical prote
15	36	80.0	424	2	conserved hypochr
16	36	80.0	472	2	omega-6 desaturase
17	36	80.0	754	2	hypothetical prote
18	36	80.0	1041	2	probable component
19	36	77.8	111	2	hypothetical prote
20	35	77.8	226	1	probable membrane
21	35	77.8	458	1	hypothetical prote
22	35	77.8	509	2	probable transemb
23	35	77.8	718	2	4-hydroxyphenylac
24	34	75.6	96	4	hypothetical prote
25	34	75.6	151	2	hypothetical prote
26	34	75.6	179	2	ribosomal protein
27	34	75.6	219	2	hypothetical prote
28	34	75.6	266	2	conserved hypochr
29	34	75.6	301	2	transcription regn

30	34	75.6	316	2	C47092	lysyl homocyst, nas
31	34	75.6	316	2	C64963	transcription regn
32	34	75.6	316	2	E05952	transcription regn
33	34	75.6	316	2	G00906	transcription regn
34	34	75.6	329	2	A01991	hypothetical prote
35	34	75.6	358	1	S41876	site-specific DNA
36	34	75.6	358	1	A97295	cellulose methyl
37	34	75.6	394	2	E04135	putative SAE DNA-b
38	34	75.6	424	2	C05891	omega-6 desaturase
39	34	75.6	492	2	E02923	alpha-1,4-glucose
40	34	75.6	497	2	T40586	nucleolar protein
41	34	75.6	522	2	E04442	nucleolar protein
42	34	75.6	559	2	E05104	hypothetical prote
43	34	75.6	799	2	T48929	serine/threonine P
44	34	75.6	910	2	C69456	subtilisin serinai
45	33.5	74.4	151	2	C03912	probable membrane

## ALIGNMENTS

### RESULT 1

T08136

probable omega-6 desaturase (F01144.99) precursor, chloroplast - rape

C1Species: Brassica napus (rape)

C1Date: 21 May 1999 #sequence\_revision 21-May-1999 #ext\_change 08 Oct 1999

C1Accession: T08136

P.Hilt, W.D., Callison, T.C., Roth, J.F., Kinney, A.C., Chou, P.L., Tatro, N.D.

Plant Physiol. 105: 635-641, 1994

A1Title: Cloning of a higher-plant plastid omega-6 fatty acid desaturase cDNA and the oxyl

A1Reference number: Z16109; MUID:94345008; PMID:8066133

A1Accession: T08136

A1Status: preliminary, translated from GB/E081/DD81

A1Molecule type: mRNA

A1Residues: 1-443 <HT>

A1Cross-reference: EMBL:L29114, NID:J01001, EMBL:M00111, EMBL:G01111

A1Experimental source: seed

C1Genetics:

C1Superfamily: omega-3 fatty acid desaturase

C1Keywords: chloroplast, fatty acid metabolism, oxidoreductase

Query Match: 88.9%, score 40; DB 2; length 443;

Best Local Similarity: 83.3%; Pred. No. 31;

Matches: 5; Conservative: 0; Mismatches: 1; Indels: 0; Gaps: 0;

QY 1 WPMWP 6

DB 252 WPMWP 257

RESULT 2

D05362

hypothetical protein AT493050 [unreviewed] - Arabidopsis thaliana

C1Species: Arabidopsis thaliana (mouse ear cress)

C1Date: 16 Feb 2001 #sequence\_revision 16 Feb 2001 #ext\_change 12 May 2002

C1Accession: D05362

P1Accession: The European Union Arabidopsis Genome Sequencing Consortium, The Cold Spring

Harbor, 100-777, 1999

A1Title: Sequence and analysis of the genome of the plant Arabidopsis thaliana.

A1Reference number: A00001, NID:000001, EMBL:000001, EMBL:000001

A1Molecule type: DNA

A1Residues: 1-448 <STO>

A1Cross-reference: EMBL:000001, NID:000001, EMBL:000001, EMBL:000001

C1Genetics:

C1Superfamily: omega-3 fatty acid desaturase

C1Keywords: chloroplast, fatty acid metabolism, oxidoreductase

Query Match: 88.9%, score 40; DB 2; length 448;

Best Local Similarity: 83.3%; Pred. No. 31;



F1453-477/Domain: cytosolic status predicted <SVS>

F132-38-36-45, 173, 176, 154, 194, 229, 266, 302, 345/sequence bonds #status experimental

F167-01-101-179, 205, 241, 254, 260, 120, 130, 170, 412/Binding site: carbohydrate (asn) (conserved)

Query Match 84.4%; Score 38; DB 1; Length 477;

Best Local Similarity 90.0%; Pred. No. 69;

Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRWH 5

DB 4 WVRWH 8

RESULT 7

A:Accession: A56853

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-822 <NA>

A:Cross-references: GR 010140, MID 060311, FIDR AAA0171.1, FID 310791

A:Reference number: 1-822 <NA>

A:Accession: A56853

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-822 <NA>

A:Cross-references: GR 010140, MID 060311, FIDR AAA0171.1, FID 310791

A:Reference number: 1-822 <NA>

A:Accession: A56853

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-822 <NA>

A:Cross-references: GR 010140, MID 060311, FIDR AAA0171.1, FID 310791

A:Reference number: 1-822 <NA>

A:Accession: A56853

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-822 <NA>

A:Cross-references: GR 010140, MID 060311, FIDR AAA0171.1, FID 310791

A:Reference number: 1-822 <NA>

A:Accession: A56853

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-822 <NA>

A:Cross-references: GR 010140, MID 060311, FIDR AAA0171.1, FID 310791

G01586

C:Species: Homo sapiens (man)

C:Date: 21-Dec-1998 #sequence\_revision 06-Jun-1997 #text\_change 20-Jun-2000

C:Accession: G01586

C:Reference number: G07954

A:Accession: G01586

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-591 <YEM>

A:Cross-references: EMBL:U08247; MID:G01224; FIDR:AAA07339.1; FID:G01224

Query Match 82.2%; Score 37; DB 2; Length 591;

Best Local Similarity 66.7%; Pred. No. 1000;

Matches 4; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 WVRWH 6

DB 350 WVRWH 355

RESULT 9

A:Accession: G01586

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: EMBL:AB010905; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: GR:AB010905; MID:G01224; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: GR:AB010905; MID:G01224; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

G01586

C:Species: Homo sapiens (man)

C:Date: 21-Dec-1998 #sequence\_revision 06-Jun-1997 #text\_change 20-Jun-2000

C:Accession: G01586

C:Reference number: G07954

A:Accession: G01586

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: mRNA

A:Residues: 1-591 <YEM>

A:Cross-references: EMBL:U08247; MID:G01224; FIDR:AAA07339.1; FID:G01224

Query Match 82.2%; Score 37; DB 2; Length 591;

Best Local Similarity 66.7%; Pred. No. 1000;

Matches 4; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 WVRWH 6

DB 350 WVRWH 355

RESULT 10

A:Accession: G01586

A:Status: preliminary; translated from GR/EMBL/DB

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: EMBL:AB010905; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: GR:AB010905; MID:G01224; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

A:Cross-references: GR:AB010905; MID:G01224; FIDR:AAA01154.1

A:Reference number: 1-195 <NA>

A:Accession: G01586

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-195 <NA>

[illegible][illegible]

Db 362 WLRWH 366

RESULT 15  
T07742

C:Species: omega-6 desaturase, chloroplast - soybean

C:Date: 14-May-1999 #sequence\_revision 14-May-1999 #text\_change 08-Oct-1999

C:Accession: T07742

R:Hit: W.D. Carlson, T.J. Roach, T.P. Kinney, A.J. Stocco, Y.L. Yip, N.S.

Plant Physiol. 105, 635-641, 1994

A:Title: Cloning of a higher plant plastid omega-6 fatty acid desaturase cDNA and its ex

A:Reference number: 216183; MIM:6445008; PMID:8066133

A:Accession: T07742

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-424 &lt;Hit&gt;

A:Cross-references: EMBL:L29215, NID:6459961, PIR: A440104.1, PIR: Q459962

A:Experimental source: seed

C:Genetics:

A:Genome: nuclear

C:Superfamily: omega-3 fatty acid desaturase

C:Keywords: chloroplast

## Query Match

80.0%; Score 36; DB 2; Length 424;

Best local similarity 66.7%; Pred. No. 1.3e+02;

Matches 4; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 WVRWHF 6

Db 251 WVRWHF 256

Search completed: February 20, 2003, 10:50:00  
Job time: 13 secs





GenScore version 5.1.3  
Copyright 1997-2003 Copyright 1993

SW protein - protein search, using SW model

February 20, 2003, 10:41:55, Search time 6.66667 seconds

(without alignment) 37 329 Million cell updates/sec

Title: US-09-912 414 2

Perfect score: 45

Sequence: 1 WWPWRF 6

Scoring table: BLOSUM62DX

Gapop 10.0, Gapext 0.5

Searched: 112892 seqs, 41476138 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: SwissProt\_40.1

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	PR	ID	Description
1	42	93.3	479	1	CATA_PSEPU	Q59714 pseudomonas
2	40	88.9	443	1	PROCT_BPAHA	P46727 brassica ca
3	40	88.9	448	1	FRAC_APAHA	P46727 arabidopsis
4	39	86.7	482	1	CATA_OXVEE	Q27312 orthococcia
5	38	84.4	822	1	TPFR_HYMAN	Q16623 hemo sapien
6	36	80.0	424	1	IFEC_SOYAN	P48628 glycine max
7	36	80.0	788	1	PCSB_XANAC	P58913 xanthomonas
8	36	80.0	1169	1	STVA_DPOME	P29194 staphylo
9	35	77.8	226	1	YPO7_METUA	Q29644 methanococ
10	35	77.8	718	1	PLSR_CAFER	Q29644 caenorhabd
11	34	75.6	148	1	PE12_METUA	P49697 methanococ
12	34	75.6	316	1	CR1_ECOLI	Q47084 escherichia
13	34	75.6	316	1	CR1_KIFAR	Q08568 klebsiella
14	34	75.6	358	1	MTG1_YANRP	Q45071 caulobacter
15	34	75.6	372	1	GP45_HUMAN	Q9Y5Y2 homo sapien
16	34	75.6	373	1	GP45_MOUSE	Q9E934 mus musculu
17	34	75.6	495	1	ACFN_ASPAT	P46849 aspergillus
18	34	75.6	799	1	AFCK_STPPO	P54741 streptomyce
19	34	75.6	807	1	AFCK_STPPO	P54741 streptomyce
20	34	75.6	1698	1	YOF6_HUMAN	Q14948 homo sapien
21	33.5	74.4	1811	1	PDRG_YEAST	Q07784 saccharomyc
22	33	73.3	744	1	YINA_CAFER	P50507 caenorhabd
23	33	73.3	376	1	MTS1_PHEME	Q10560 thirubium m
24	33	73.3	377	1	MTS1_PHEME	Q10560 thirubium m
25	33	73.3	366	1	PCP_ECOLI	P39247 escherichia
26	33	73.3	439	1	YAL2_ARATH	Q04656 arabidopsis
27	33	73.3	478	1	CATA_LACSR	P33569 streptomyce
28	33	73.3	483	1	BCA_STRVL	P24422 micrococcu
29	33	73.3	497	1	CATA_MICLU	Q9XZ45 toxoplasma
30	33	73.3	502	1	CATA_TOXOP	Q14460 saccharomyc
31	33	73.3	504	1	SIR1_YEAST	Q12499 saccharomyc
32	33	73.3	511	1	NOPS_YEAST	Q9Y2X1 homo sapien
33	33	73.3	529	1	NOPS_HUMAN	

34	33	73.3	534	1	NOPS_PAT	Q9Q284 ratius norv
35	33	73.3	614	1	WQ3E_YEAST	P53293 saccharomyc
36	33	73.3	822	1	TPFR_MOUSE	P12309 mus musculu
37	33	73.3	821	1	TPFR_PAT	Q46044 ratius norv
38	33	73.3	1015	1	FDNG_ECOLI	P24183 escherichia
39	33	73.3	1014	1	FDNG_ECOLI	P24183 escherichia
40	32	71.1	129	1	YHME_YEAST	P38857 saccharomyc
41	32	71.1	146	1	YHME_YEAST	P38857 saccharomyc
42	32	71.1	287	1	YHME_PHEME	P77205 escherichia
43	32	71.1	386	1	PL4_DRECA	P49165 urechis cau
44	32	71.1	395	1	DUTR_BACSU	P39580 bacillus su
45	32	71.1	396	1	RVA_XENMA	P03127 xenopus lae

## ALIGNMENTS

RESULT 1  
CATA\_PSEPU STANDARD; PRT: 479 AA.  
ID CATA\_PSEPU  
AC Q59714:  
DT 30-MAY-2000 (rel. 39, Created)  
DT 30-MAY-2000 (rel. 39, Last sequence update)  
DT 30 MAY 2000 (rel. 39, Last annotation update)  
DE Catalase (EC 1.11.1.6).  
GN KATA OF CATA.  
OS Pseudomonas putida.  
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;  
OC Pseudomonas.  
OX NCBI\_TaxID=303;  
RN (1)  
RP SEQUENCE FROM N.A.  
RC STRAIN=corvallis;  
RX MERRILL:09010901; PubMed:01090901;  
PA Kim Y C, Miller C P, Anderson A U;  
RT Identification of adjacent genes encoding the major catalase and a  
RT bacterioferritin from the plant-beneficial bacterium pseudomonas  
RT putida.";  
RL Gene 199:219-224(1997).  
CC FUNCTION: CATALASE: HYDROGEN PEROXIDE IN WATER AND OXYGEN, FORMS  
CC TO PROTECT CELLS FROM THE TOXIC EFFECTS OF HYDROGEN PEROXIDE.  
CC -1- CATALYTIC ACTIVITY: 2 H2O2(O2) -> 2 H2O.  
CC -1- COFACTOR: HEME GROUP.  
CC -1- ENZYME REGULATION: ACTIVATED BY PEROXIDE.  
CC -1- SIMILARITY: BELONGS TO THE CATALASE FAMILY.  
CC This SWISS-PROT entry is copyright. It is provided through a collaboration  
CC between the Swiss Institute of Bioinformatics and the EMBL databank.  
CC the European Bioinformatics Institute. There are no restrictions on its  
CC use by non profit institutions as long as its content is in no way  
CC modified and this statement is not removed. Usage by and for commercial  
CC entities requires a license agreement (See <http://www.isb-eb.ch/announcements/>  
CC or send an email to [license@isb.ch](mailto:license@isb.ch)).  
DR EMBL: U63511; AAB88219.1; ..  
DR HSSP: F42311; 2CAE.  
DR InterPro: IPR002226; Catalase.  
DR Pfam: PF00199; catalase.1.  
DR PRINTS: PR00067; CATALASE.  
DR PRODOM: PD000510; Catalase.1.  
DR PROSITE: PS00437; CATALASE.1; 1.  
DR PROSITE: PS00438; CATALASE.2; 1.  
FW Oxidoreductase; peroxidase; iron, heme; hydrogen peroxide.  
FT ACT SITE 53  
FT ACT SITE 53  
FT ACT SITE 126  
FT BINDING 336 336 PROXIMAL HEME LIGAND (BY SIMILARITY).  
SQ SEQUENCE 479 AA: 53381 MW: 115300.6778571 CRC64;  
Query Match 93.3%, Score 42, DE 1, Length 479;  
Best local similarity 83.3%, Pred. No. 3.4;  
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
QY 1 WWPWRF 6



CC 1 FUNCTION: THE PROLACTIN-MEMBRANE FATTY ACID DESATURASE INTERFERES  
 CC 2 WITH THE SECOND INTERFERED IN THE BIOSYNTHESIS OF 16:3 AND 18:3 FATTY  
 CC 3 ACIDS. INTERFERED INTERFERES IN THE BIOSYNTHESIS OF PLANT MEMBRANES. IT IS THOUGHT  
 CC 4 TO USE PEROXYDIN AS AN ELECTRON DONOR AND TO ACT ON FATTY ACIDS  
 CC 5 ESTERIFIED TO Glycerolipids, Sphingolipids AND PHOSPHATIDYLCHOLINE.  
 CC 6 ESTERIFIED TO Glycerolipids, Sphingolipids AND PHOSPHATIDYLCHOLINE.  
 CC 7 SUPPLEMENTAL LOCATION: CHLOROPLAST, MEMBRANE-BINDING (PDB-ENTRY)  
 CC 8 DEVELOPMENTAL STAGE: HIGHEST LEVELS FOUND IN EXPANDING LEAVES.  
 CC 9 DOMAIN: THE HISTIDINE BOX DOMAINS MAY CONTAIN THE ACTIVE SITE  
 CC 10 AND 15 PER INVOLVED IN METAL ION BINDING.  
 CC 11 SIMILARITY: BELONGS TO THE FATTY ACID DESATURASE FAMILY  
 CC 12  
 CC 13 This SWISS-PROT entry is copyright. It is produced through a collaboration  
 CC 14 between the Swiss Institute of Bioinformatics and the EMBL, contribution  
 CC 15 the European Bioinformatics Institute. There are no restrictions on its  
 CC 16 use by non-profit institutions as long as its content is in no way  
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 CC 19 or send an email to [license@sib-sib.ch](mailto:license@sib-sib.ch))  
 CC 20  
 CC 21 EMBL: U09503; AAA92800.1;  
 CC 22 EMBL: A1023108; CAA91908.1;  
 CC 23 EMBL: A161578; CAB98813.1;  
 CC 24 EMBL: AY045673; AAY77979.1;  
 CC 25 EMBL: AY045672; AAI24196.1;  
 CC 26 EMBL: AY045652; AAI24340.1;  
 CC 27 InterPro: IPR001225; FA\_desaturase.  
 CC 28 Pfam: PF00487; FA\_desaturase; 1  
 CC 29 ProDom: P000081; FA\_desaturase; 1  
 CC 30 Oxytruncatella; Fatty acid biosynthesis; Chloroplast; Membrane;  
 CC 31 Transmembrane protein;  
 CC 32 TRANSIT 1 69 CHLOROPLAST (BY SIMILARITY);  
 CC 33 FT CHAIN 70 448 OMEGA-6 FATTY ACID DESATURASE;  
 CC 34 FT DOMAIN 171 175 HISTIDINE BOX-1;  
 CC 35 FT DOMAIN 207 211 HISTIDINE BOX-2;  
 CC 36 FT DOMAIN 367 371 HISTIDINE BOX-3;  
 CC 37 SEQUENCE 448 AA; 5175 MW; 33477.56 kDa; 67.2% GC;  
 CC 38  
 CC 39 Query Match 88.9% Score 40; PB 1; Length 448;  
 CC 40 Best Local Similarity 93.3%; Freq. No. 5.3;  
 CC 41 Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 CC 42  
 CC 43 QY 1 WVRWHF 6  
 CC 44 DB 257 WVRWHF 262  
 CC 45  
 CC 46 RESULT 4  
 CC 47 DATA ONCE STANDARD; FRT; 482 AA;  
 CC 48 ID CATA\_ONCE  
 CC 49 AC Q27710; Q85499;  
 CC 50 DT 30-MAY-2000 (Rel. 39, Created)  
 CC 51 DT 30-MAY-2000 (Rel. 39, Last sequence update)  
 CC 52 DT 10-MAY-2000 (Rel. 39, Last annotation update)  
 CC 53 DE Catalase (EC 1.11.1.6)  
 CC 54 GN CAT  
 CC 55 OS *Onchocerca volvulus* (nematode)  
 CC 56 NC Bacteria; Proteobacteria; alpha subdivision; Rickettsiales;  
 CC 57 CC Rickettsiales; Wolbachiales  
 CC 58 NCBI\_TaxID=77551;  
 CC 59 RN [1]  
 CC 60 RP SEQUENCE FROM N.A.  
 CC 61 RA Henkels-Duetsch K.J.; Walter V.H.O.; Wildenburg G.; Blaxter M.;  
 CC 62 Walter R.D.;  
 CC 63 "Gene structure, activity and localization of a catalase from  
 CC 64 intracellular bacteria in *Onchocerca volvulus*,"  
 CC 65 submitted (MAY-1998) to the EMBL/Genbank/DBP databases.  
 CC 66 CC EMBL: U09503; AAA92800.1;  
 CC 67 CC EMBL: U09503; AAA92800.1;  
 CC 68 CC EMBL: U09503; AAA92800.1;  
 CC 69 CC EMBL: U09503; AAA92800.1;  
 CC 70 CC EMBL: U09503; AAA92800.1;  
 CC 71 CC EMBL: U09503; AAA92800.1;  
 CC 72 CC EMBL: U09503; AAA92800.1;  
 CC 73 CC EMBL: U09503; AAA92800.1;  
 CC 74 CC EMBL: U09503; AAA92800.1;  
 CC 75 CC EMBL: U09503; AAA92800.1;  
 CC 76 CC EMBL: U09503; AAA92800.1;  
 CC 77 CC EMBL: U09503; AAA92800.1;  
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 CC 80 CC EMBL: U09503; AAA92800.1;  
 CC 81 CC EMBL: U09503; AAA92800.1;  
 CC 82 CC EMBL: U09503; AAA92800.1;  
 CC 83 CC EMBL: U09503; AAA92800.1;  
 CC 84 CC EMBL: U09503; AAA92800.1;  
 CC 85 CC EMBL: U09503; AAA92800.1;  
 CC 86 CC EMBL: U09503; AAA92800.1;  
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 CC 89 CC EMBL: U09503; AAA92800.1;  
 CC 90 CC EMBL: U09503; AAA92800.1;  
 CC 91 CC EMBL: U09503; AAA92800.1;  
 CC 92 CC EMBL: U09503; AAA92800.1;  
 CC 93 CC EMBL: U09503; AAA92800.1;  
 CC 94 CC EMBL: U09503; AAA92800.1;  
 CC 95 CC EMBL: U09503; AAA92800.1;  
 CC 96 CC EMBL: U09503; AAA92800.1;  
 CC 97 CC EMBL: U09503; AAA92800.1;  
 CC 98 CC EMBL: U09503; AAA92800.1;  
 CC 99 CC EMBL: U09503; AAA92800.1;  
 CC 100 CC EMBL: U09503; AAA92800.1;

CC 1 CATALYTIC ACTIVITY: 2 H<sub>2</sub>O<sub>2</sub> + 2 H<sub>2</sub>O  
 CC 2 - COCAINE: HEME GROUP  
 CC 3 - SIMILARITY: BELONGS TO THE CATALASE FAMILY  
 CC 4  
 CC 5 This SWISS-PROT entry is copyright. It is produced through a collaboration  
 CC 6 between the Swiss Institute of Bioinformatics and the EMBL, contribution  
 CC 7 the European Bioinformatics Institute. There are no restrictions on its  
 CC 8 use by non-profit institutions as long as its content is in no way  
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 CC 11 or send an email to [license@sib-sib.ch](mailto:license@sib-sib.ch))  
 CC 12  
 CC 13 EMBL: X82176; CAA57666.1;  
 CC 14 EMBL: AF060070; AAC70431.1;  
 CC 15 EMBL: P42321; CAAE  
 CC 16 InterPro: IPR002226; Catalase.  
 CC 17 Pfam: PF00199; Catalase.  
 CC 18 PRINTS: PR00667; CATALASE  
 CC 19 ProDom: P000081; Catalase; 1  
 CC 20 PROSITE: PS00437; CATALASE.1;  
 CC 21 PROSITE: PS00439; CATALASE.2;  
 CC 22 Oxytruncatella; Peroxidase; Int. Heme; Hydrogen peroxide;  
 CC 23 ACT SITE 55 55  
 CC 24 ACT SITE 128 128  
 CC 25 BINDING 338 338  
 CC 26 SEQUENCE 482 AA; 53700 MW; 33477.56 kDa; 67.2% GC;  
 CC 27  
 CC 28 Query Match 86.7% Score 19; PB 1; Length 482;  
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 CC 30 Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC 31  
 CC 32 QY 1 WVRWHF 6  
 CC 33 DB 211 WVRWHF 216  
 CC 34  
 CC 35 RESULT 5  
 CC 36 TRKB HUMAN  
 CC 37 ID TRKB\_HUMAN STANDARD; FRT; 422 AA;  
 CC 38 AC Q27710; Q85499;  
 CC 39 DT 01-NOV-1997 (Rel. 35, Created)  
 CC 40 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 CC 41 DT 15-JUN-2002 (Rel. 41, Last annotation update)  
 CC 42 DE BDNF/NT-3 growth factors receptor precursor (FC 2 7 1 112) (TrkB)  
 CC 43 DE Tyrosine kinase (GPI45-TrkB) (TrkB-B)  
 CC 44 GN NTRK2; OP TrkB  
 CC 45 OS Homo sapiens (Human)  
 CC 46 NC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 CC 47 CC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo;  
 CC 48 NCBI\_TaxID=9606;  
 CC 49 RN [1]  
 CC 50 RP SEQUENCE FROM N.A. (LONG FORM)  
 CC 51 RA Nakagawa A.; Liu X.-G.; Ikegaki N.; White P.S.; Yamashiro D.J.;  
 CC 52 Nakagawa A.; Liu X.-G.; Ikegaki N.; White P.S.; Yamashiro D.J.;  
 CC 53 "Cloning and chromosomal localization of the human TrkB P tyrosine  
 CC 54 kinase receptor gene (NTRK2)."  
 CC 55 Genomics 25:538-546(1995).  
 CC 56 RN [2]  
 CC 57 RP SEQUENCE FROM N.A. AND PARTIAL SEQUENCE (LONG AND TRUNCATED FORMS).  
 CC 58 RA MEDLINE:9523473; PubMed:7823156;  
 CC 59 RA Shellen D.L.; Sutherland V.; Grupp J.; Camerato T.; Armanini M.P.;  
 CC 60 Phillips H.S.; Carroll K.; Spencer S.D.; Levinson A.D.;  
 CC 61 "Human TrkB molecular cloning, tissue distribution, and expression  
 CC 62 of extracellular domain immunoadhesins."  
 CC 63 J. Neurosci. 15:477-491(1995)  
 CC 64 RN [3]  
 CC 65 RP SEQUENCE FROM N.A. (TRUNCATED FORMS).  
 CC 66 RA MEDLINE:9523473; PubMed:7823156;  
 CC 67 CC MEDLINE:9523473; PubMed:7823156;  
 CC 68 CC MEDLINE:9523473; PubMed:7823156;  
 CC 69 CC MEDLINE:9523473; PubMed:7823156;  
 CC 70 CC MEDLINE:9523473; PubMed:7823156;  
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 CC 72 CC MEDLINE:9523473; PubMed:7823156;  
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 CC 98 CC MEDLINE:9523473; PubMed:7823156;  
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 CC 100 CC MEDLINE:9523473; PubMed:7823156;









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RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 99156852; PubMed 10036181;
RA Marchese A., Sawatzko M., Nguyen T., Cheng R., Heng H.H., Nowak T.,
RA Im D.-S., Lynch K.P., George S.P., O'Dowd B.F.;
RL "Discovery of three novel orphan G-protein-coupled receptors.";
RL Genomics 56:12-21(1999).
CC -!- FUNCTION: Orphan receptor. May play a role in brain function.
CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Expressed in brain; detected in the basal
CC -!- TISSUE SPECIFICITY: Expressed in brain; detected in the basal
CC -!- FORTHRAIN, frontal cortex, and caudate, but not in thalamus,
CC hippocampus, or putamen.
CC -!- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
CC
CC This Swiss-Prot entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC or send an email to license@sib-sib.ch).
CC
DR EMBL: AF18266; AAD2106.1; -
DR Genew: HNC4503; GPR45.
DR MIM: 604838; -
DR InterPro: IPR00276; GPCR_Phodspn.
DR Pfam: PF00001; 7tm_1; 1.
DR PRINTS: FRC0237; GPCR_HUMAN.
DR PROSITE: PS00276; G-PROTEIN_RECEP_F1_1; FALSE_NEG.
DR PROSITE: PS00762; G-PROTEIN_RECEP_F1_2; 1.
KW G-protein coupled receptor, transmembrane, glycoprotein,
FT DOMAIN 1 38
FT TRANSMEM 39 59
FT DOMAIN 60 75
FT TRANSMEM 76 96
FT DOMAIN 97 109
FT TRANSMEM 110 130
FT DOMAIN 131 149
FT TRANSMEM 150 170
FT DOMAIN 171 198
FT TRANSMEM 199 219
FT DOMAIN 220 268
FT TRANSMEM 269 289
FT DOMAIN 290 305
FT TRANSMEM 306 326
FT DOMAIN 327 372
FT CARBOHYD 4 4
FT CARBOHYD 17 17
FT CARBOHYD 29 29
FT SEQUENCE 372 AA; 47000 MW; R4R4F7AEDE20199 CPG64;

Query Match 75.6%; Score 34; DB 1; Length 372;
Best Local Similarity 70.0%; Pred. No. 57;
Matches 5; Conservation 0; Mismatches 0; Indels 0; Gaps 0;
OY 2 VRWHF 6
DB 98 VRWHF 102

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Search completed: February 20, 2003, 10:48:01  
 Job time : 8.66667 secs











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ID 08WXJ4 PRELIMINARY: PRT: 70 AA.
AC 08WXJ4:
DT 01-MAR-2002 (TEMBLrel_20, Created)
DT 01-MAR-2002 (TEMBLrel_20, Last sequence update)
DT 01-MAR-2002 (TEMBLrel_20, Last annotation update)
DE Neurite growth promoting tyrosine kinase type 2 (Fragment).
GN NTRK2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE 2165693; PubMed 11798192;
RA Stojilov P., Gastien E., Stamm S.;
RT "Analysis of the Human TRK Gene Genomic Organization Reveals Novel
RT Tyk Isoforms, Unusual Gene Length, and Splicing Mechanism."
RL Biochem Biophys Res Commun 300:1064-1067(2002).
RL EMBL: AF410902; AAL67966.1;
DR InterPro: IPR000372; IPR_Nterm
DR Pfam: PF01462; LRRNT_1;
DR SMART: SM00013; LRRNT_1;
FW Receptor: Kinase.
FT NON_TER
FT NON_TER
SQ SEQUENCE 70 AA; 344 MW; 18984040000/23A CPC64;

Query Match 84.4%; Score 38; DB 4; Length 70;
Best Local Similarity 80.0%; Pred No. 22;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRWH 5
Db 4 WVRWH 8

RESULT 14
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ID 095K65
AC 095K65:
DT 01-DEC-2001 (TEMBLrel_19, Created)
DT 01-FEB-2001 (TEMBLrel_19, Last sequence update)
DT 01-JUN-2002 (TEMBLrel_21, Last annotation update)
DE Hypothetical 20.3 kDa protein.
DR Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=MEDULLA OBLONGATA;
RA Oseada N., Hida M., Kiyoda T., Tanuma P., Iseki K., Hirai M., Terao K.,
RA Suzuki Y., Sugano S., Hashimoto K.;
RT "Isolation of full-length cDNA clones from macaque brain cDNA
RT libraries."
RL Submitted (JUL-2001) to the EMBL/GenBank/DDBJ databases.
RL EMBL: AB066522; BAB62201.1;
DR InterPro: IPR000372; LRR_Nterm.
DR Pfam: PF01462; LRRNT_1;
KM Hypothetical protein.
SQ SEQUENCE 178 AA; 20233 MW; A7800A4355F058F CPC64;

Query Match 84.4%; Score 38; DB 6; Length 178;
Best Local Similarity 80.0%; Pred No. 55;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRWH 5
Db 4 WVRWH 8

RESULT 15

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08WXJ6
ID 08WXJ6 PRELIMINARY: PRT: 537 AA.
AC 08WXJ6:
DT 01-MAR-2002 (TEMBLrel_20, Created)
DT 01-MAR-2002 (TEMBLrel_20, Last sequence update)
DT 01-MAR-2002 (TEMBLrel_20, Last annotation update)
DE Neurite growth promoting tyrosine kinase type 2 (Fragment).
GN NTRK2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE 2165693; PubMed 11798192;
RA Stojilov P., Gastien E., Stamm S.;
RT "Analysis of the Human TRK Gene Genomic Organization Reveals Novel
RT Tyk Isoforms, Unusual Gene Length, and Splicing Mechanism."
RL Biochem Biophys Res Commun 300:1064-1067(2002).
RL EMBL: AF410902; AAL67966.1;
DR InterPro: IPR000372; IPR_Nterm
DR Pfam: PF00047; IG_1;
DR Pfam: PF00560; LRR_1;
DR Pfam: PF01462; LRRNT_1;
DR Pfam: PF01462; LRRNT_1;
DR SMART: SM00409; IG_1;
DR SMART: SM00408; IG02_1;
DR SMART: SM00682; LRRCT_1;
DR SMART: SM00013; LRRNT_1;
FW Receptor: Kinase.
FT NON_TER
SQ SEQUENCE 537 AA; 59166 MW; 5A8FA252A38710C1 CPC64;

Query Match 84.4%; Score 38; DB 4; Length 537;
Best Local Similarity 80.0%; Pred No. 16602;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 WVRWH 5
Db 4 WVRWH 8

Search completed: February 20, 2003, 10:43:18
Job time: 25.3333 secs

```





GenCore version 5.1.3  
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OW protein - protein search, using sw mode

Run on: February 20, 2003, 11:24:02 / Search time 12 seconds  
(without alignments)  
48,067 Million cell updates/sec

Title: US-09 912 414 9

Perfect score: 6

Sequence: 1 WXXWXP 6

Scoring table: oligo

Gapop 60.0, Gapext 60.0

Searched: 283224 seqs, 96134422 residues

Word size: 0

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summations

Database: PIP\_731  
1: p1r1  
2: p1r2  
3: p1r3  
4: p1r4

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the best hit found, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	6	100.0	21	2	PH1731
2	6	100.0	49	2	I46641
3	6	100.0	59	2	E82844
4	6	100.0	41	2	H91106
5	6	100.0	65	2	S23164
6	6	100.0	71	2	F84168
7	6	100.0	73	2	AH7288
8	6	100.0	76	2	T14724
9	6	100.0	78	2	S94443
10	6	100.0	79	2	M47760
11	6	100.0	83	2	A97568
12	6	100.0	86	2	T42185
13	6	100.0	87	2	H86752
14	6	100.0	92	2	C70609
15	6	100.0	92	2	I01627
16	6	100.0	93	2	G87098
17	6	100.0	97	2	E84147
18	6	100.0	99	2	C84022
19	6	100.0	101	2	C71519
20	6	100.0	103	2	F82266
21	6	100.0	105	2	F80184
22	6	100.0	106	2	I64171
23	6	100.0	106	2	C82715
24	6	100.0	108	2	C72913
25	6	100.0	109	2	A91164
26	6	100.0	109	2	C82241
27	6	100.0	109	2	A92505
28	6	100.0	109	2	F81057
29	6	100.0	109	2	A12029

30	6	100.0	110	2	AD0127	conserved hypothet
31	6	100.0	110	2	AG1523	hypothetical prote
32	6	100.0	116	2	P00066	13 heavy chain pro
33	6	100.0	116	2	S82388	13 heavy chain pro
34	6	100.0	117	2	A95381	hypothetical prote
35	6	100.0	118	2	S24527	13 heavy chain V r
36	6	100.0	119	2	S24522	13 heavy chain V r
37	6	100.0	119	2	S24492	13 heavy chain V r
38	6	100.0	119	2	S24513	13 heavy chain V r
39	6	100.0	119	2	S24493	13 heavy chain V r
40	6	100.0	119	2	S24491	13 heavy chain V r
41	6	100.0	119	2	S24525	13 heavy chain V r
42	6	100.0	119	2	S24520	13 heavy chain V r
43	6	100.0	119	2	S24518	13 heavy chain V r
44	6	100.0	119	2	S24511	13 heavy chain V r
45	6	100.0	119	2	S24494	13 heavy chain V r

#### ALIGNMENTS

##### RESULT 1

PH1731

Ig heavy chain V region (clone GCC-11) - mouse (fragment)

C.Species: Mus musculus (house mouse)

C.Date: 24 Feb 1994 #Seq-ident: 100.00% #Align: 24 Feb 1994 #Gap: 0.00% #Align: 24 Feb 1994

C.Accession: PH1731

P.McMurry, Williams, M.G., Melton, M.J., Lator, F.A., Hessel, G.J.V.

J. Exp. Med. 178, 245-307, 1993

A>Title: Antigen-driven B cell differentiation in vivo.

A.Reference number: PH1675; PMID:93301607; PMID:815385

A.Accession: PH1731

A.Molecule type: mRNA

A.Pesides: 1-21 <MCH>

A.Experimental source: B cell

A.Note: The authors translated the cDNA ACA for residue 13 as Ala

Superficially, immunoglobulin V region, (residue 13) is 13-37

C.Keywords: heterotrimer; immunoglobulin

Query Match: 100.0%; Score 6; PR 2; Length 21;

Best Local Similarity: 50.0%; Pred. No. 146641; Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6

DB 16 WXSWSF 21

##### RESULT 2

I46641

185 kDa glycoprotein - mouse (fragment)

C.Species: Mus musculus (house mouse)

C.Date: 22 Jul 1994 #Sequence\_ident: 100.00% #Align: 22 Jul 1994

C.Accession: I46641

A.Status: preliminary; translated from cDNA/EMBL/GenBank

A.Molecule type: DNA

A.Pesides: 1-49 <PE>

A.Reference number: EMBL X00004; MIM 312100; F000000000; F000000000

A.Creation: 1994

A.Keywords: ATP; phosphoprotein

Query Match: 100.0%; Score 6; PR 2; Length 49;

Best Local Similarity: 50.0%; Pred. No. 146641; Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6

DB 16 WXSWSF 21



A:Accession: AH2788  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-73 <KUR>  
 A:Cross-references: GR A600498, F1FN AA42726.1, F1D J1741165, GCESE GN0186  
 A:Experimental source: strain C8A (Dupont)  
 C:Genetics:  
 A:Gene: Atul727  
 A:Map position: circular chromosome

Query Match 100.0%; Score 6; DB 2; Length 73;  
 Best Local Similarity 50.0%; Pred No 1 9e+03;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 [...] |  
 DB 23 MWLWF 18

RESULT 8  
 T14724  
 hypothetical protein Y1091 - Yersinia pestis plasmid pMT1

C:Species: Yersinia pestis  
 C:Date: 10-Sep-1993 #sequence\_revision 30-Sep-1993 #ext\_change 04-Sep-1993  
 C:Accession: T14724; T15006  
 R:Hu, F.; Elliott, J.; McCready, P.; Skowronski, E.; Tarras, P.; Kobayashi, A.; et al.,  
 submitted to the EMBL Data Library, March 1998  
 A:Description: Structural organization of virulence determinants in three Yersinia pestis  
 A:Reference number: Y18168  
 A:Accession: T14724  
 A:Status: preliminary; translated from GR/EMBL/GENB  
 A:Molecule type: DNA  
 A:Residues: 1-76 <HUP>  
 A:Cross-references: EMBL A093447, F1D G290096, F1D G290267, F1D AAC11243.1  
 R:Rindler, L.E.; Plano, G.V.; Portland, V.; Mayhew, G.F.; Plattner, F.P.  
 Infect. Immun. 66, 5731-5742, 1998  
 A:Title: Complete DNA sequence and detailed analysis of the Yersinia pestis KIMS plasmid  
 A:Reference number: Z18268; M1D:09043898; PMID:9826348  
 A:Accession: T15006  
 A:Status: preliminary; translated from GR/EMBL/GENB  
 A:Molecule type: DNA  
 A:Residues: 1-76 <HUP>  
 A:Cross-references: EMBL A093447, F1D G290096, F1D G290267, F1D AAC11243.1  
 C:Genetics:  
 A:Gene: Y1091  
 A:Genome: plasmid pMT1

Query Match 100.0%; Score 6; DB 2; Length 76;  
 Best Local Similarity 50.0%; Pred No 1 9e+03;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 [...] |  
 DB 50 WRTWNP 55

RESULT 9

S08449  
 hypothetical protein 9 - Spiroplasma virus 1  
 C:Species: Spiroplasma virus 1, SPV1  
 C:Date: 10-Jun-1991 #sequence\_revision 30-Jun-1991 #ext\_change 07-Dec-1990  
 C:Accession: S08449  
 R:Penauhin, J.; Aulio, P.; Vignault, J.C.; Bove, J.M.  
 Nucleic Acids Res. 18, 1293, 1990  
 A:Title: Complete nucleotide sequence of the genome of Spiroplasma citri virus SPV1-RNA  
 A:Reference number: S08447; M1D:0906799; PMID:2326423  
 A:Accession: S08449  
 A:Status: translation not shown  
 A:Molecule type: RNA  
 A:Residues: 1-78 <REN>  
 A:Cross-references: EMBL X1114, M1D J1700, F1D CA4727.1, F1D G08664  
 C:Genetics:  
 A:Genetic code: SGR1

Query Match 100.0%; Score 6; DB 2; Length 78;  
 Best Local Similarity 50.0%; Pred No 2e+03;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 [...] |  
 DB 42 MWLWF 47

RESULT 10

A64750  
 hypothetical protein b0249 Escherichia coli (strain K12)  
 C:Species: Escherichia coli  
 C:Date: 12-Sep-1997 #sequence\_revision 10-Sep-1997 #ext\_change 01-Mar-2002  
 C:Accession: A64750  
 R:Blattner, F.P.; Plunkett III, G.; Blythe, C.A.; Perna, N.T.; Burland, V.; Riley, M.; et al.  
 A: Rose, D.N.; Mau, B.; Shao, Y.  
 Science 277, 1453-1462, 1997  
 A:Title: The complete genome sequence of Escherichia coli K-12.  
 A:Reference number: A64750; M1D:0742617; PMID:9298603  
 A:Accession: A64750  
 A:Status: nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-79 <BLAT>  
 A:Cross-references: GR A600117, F1D G290096, F1D G290267, F1D AAC11243.1, F1D J1741165  
 A:Experimental source: strain K-12, substrain MO155

Query Match 100.0%; Score 6; DB 2; Length 79;  
 Best Local Similarity 50.0%; Pred No 2e+03;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 [...] |  
 DB 50 WRTWNP 55

RESULT 11

A97568  
 hypothetical protein AGR\_C\_3173 (imported) - Agrobacterium tumefaciens strain 289, T100  
 C:Species: Agrobacterium tumefaciens  
 C:Date: 20-Sep-2001 #sequence\_revision 30-Sep-2001 #ext\_change 11-Mar-2002  
 C:Accession: A97568  
 R:Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Planchard, M.; Gurello, B.; Goldman,  
 A.; Liu, F.; Wollam, C.; Allinger, M.; Dougherty, D.; Scott, C.; Lappas, C.; Markovitz, B.;  
 Science 294, 2303-2328, 2001  
 A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tumefaciens  
 A:Reference number: A97569; PMID:11745194  
 A:Accession: A97568  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-83 <KUR>  
 A:Cross-references: GR A600498, F1FN AA42726.1, F1D J1741165, GCESE GN0186  
 C:Genetics:  
 A:Gene: AGR\_C\_3173  
 A:Map position: circular chromosome

Query Match 100.0%; Score 6; DB 2; Length 83;  
 Best Local Similarity 50.0%; Pred No 2 1e+03;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 [...] |  
 DB 23 MWLWF 28

RESULT 12

T42185  
 conserved hypothetical protein U7085 - Escherichia coli plasmid p0157  
 C:Species: Escherichia coli  
 C:Date: 03-Dec-1990 #sequence\_revision 03-Dec-1990 #ext\_change 02-Dec-1999  
 C:Accession: T42185  
 R:Portland, V.; Shao, Y.; Perna, N.T.; Plunkett, G.; Blythe, C.A.; Riley, M.; et al.



GenCode version 5.1  
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OM protein - protein search, using sw method

Run on: February 20, 2003, 11:20:17 (Search time 11:20:17 seconds)  
(without alignments)  
21,958 Million cell updates/sec

Title: US-09-912-414-9

Perfect score: 6

Sequence: 1 WXXWKF 6

Scoring table: OLLIGOX

Gapd 60.0, Gapext 60.0

Searched: 112892 seqs, 41476328 residues

Word size: 0

Total number of hits satisfying chosen parameters: 112892

Minimum PR seq length: 0

Maximum PR seq length: 200000000

Post processing: listing first 45 summaries

Database: SwissProt\_40.1

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No	Score	Query Match	Length	PR	ID	Description
1	6	100.0	55	1	LHA2_ECTHL	Protein ectothiorhodopsin
2	6	100.0	78	1	W99_SEVIR	Protein septin
3	6	100.0	79	1	YKRF_ECOLI	Protein septin
4	6	100.0	84	1	ACHA_CROPS	Protein septin
5	6	100.0	84	1	ACHA_PELTA	Protein septin
6	6	100.0	90	1	YKRF_ECOLI	Protein septin
7	6	100.0	101	1	YKRF_MYCN	Protein septin
8	6	100.0	105	1	NICW_HUMAN	Protein septin
9	6	100.0	105	1	YKRF_HUMAN	Protein septin
10	6	100.0	108	1	NICW_HUMAN	Protein septin
11	6	100.0	116	1	HVNS_CARPAT	Protein septin
12	6	100.0	128	1	MERT_STANT	Protein septin
13	6	100.0	129	1	KVHL_HUMAN	Protein septin
14	6	100.0	129	1	YKRF_YEAST	Protein septin
15	6	100.0	138	1	YKRF_MYCN	Protein septin
16	6	100.0	139	1	YKRF_MYCN	Protein septin
17	6	100.0	145	1	HVNS_MOUSE	Protein septin
18	6	100.0	145	1	PL32_ABPPE	Protein septin
19	6	100.0	145	1	VKSF_YEAST	Protein septin
20	6	100.0	147	1	VKSF_YEAST	Protein septin
21	6	100.0	155	1	HVNS_ECOLI	Protein septin
22	6	100.0	157	1	YKRF_MYCN	Protein septin
23	6	100.0	164	1	LSPA_ECOLI	Protein septin
24	6	100.0	165	1	NSAR_VIBAL	Protein septin
25	6	100.0	165	1	LSPA_ENTAE	Protein septin
26	6	100.0	165	1	YKRF_PHOSH	Protein septin
27	6	100.0	166	1	YKRF_SEVIR	Protein septin
28	6	100.0	170	1	LSPA_PSEPI	Protein septin
29	6	100.0	171	1	YKRF_MYCN	Protein septin
30	6	100.0	176	1	VPI_BPP2	Protein septin
31	6	100.0	177	1	YKRF_HUMAN	Protein septin
32	6	100.0	182	1	VG37_BPMU	Protein septin
33	6	100.0	182	1	VG37_BPMU	Protein septin

## ALIGNMENTS

34	6	100.0	196	1	YKRF_YEAST	P46285 saccharomyces
35	6	100.0	206	1	YKRF_MYCN	P34602 escherichia
36	6	100.0	167	1	YKRF_ECOLI	P34602 escherichia
37	6	100.0	208	1	YKRF_ECOLI	P34602 escherichia
38	6	100.0	210	1	SSH5_YEAST	Q03446 saccharomyces
39	6	100.0	212	1	YKRF_YEAST	P48597 xenopus laevis
40	6	100.0	215	1	YKRF_YEAST	Q77210 aplysia californica
41	6	100.0	217	1	YKRF_HUMAN	P07720 human septin
42	6	100.0	217	1	YKRF_MYCN	P20415 mus musculus
43	6	100.0	217	1	YKRF_PSEPI	P04328 cytrichlamydomonas
44	6	100.0	219	1	ENGR_MENJA	Q57768 methanocaldococcus
45	6	100.0	220	1	P208_PSEPI	P03048 bacteriophage

RESULT 1  
LHA2\_ECTHL STANDARD; PRT; 65 AA.

AC P80103;  
DT 01-FEB-1994 (Rel. 28, Created)  
DT 01-FEB-1994 (Rel. 28, Last sequence update)  
DE Light-harvesting protein 8800/810/1020, alpha-2 chain (ENS-alpha 2)  
DE Antenna pigment protein, alpha-2 chain)  
OS Ectothiorhodospira halochloris.  
OC Bacteria, Proteobacteria, gamma subdivision, Ectothiorhodospiraceae  
OX NCBI\_TaxID:1052;  
RN (1)  
RP SEQUENCE.  
PX STPAIN-DSM 1059;  
PY MERLIN-2224936; PubMed:1577009;  
RA Wagner-Huber P., Brunstolz P.A., Bisig I., Frank G., Suter P., Zuber H.;  
RT "The primary structure of the antenna polypeptides of Ectothiorhodospira halochloris and Ectothiorhodospira halophila. Four core-type antenna polypeptides in E. halochloris and E. halophila." Eur. J. Biochem. 205:917-925(1992)  
RL  
CC "FUNCTION: ANTENNA COMPLEXES ARE LIGHT-HARVESTING SYSTEMS, WHICH TRANSMIT THE EXCITATION ENERGY TO THE REACTION CENTERS. SUBUNIT: THE CORE COMPLEX IS FORMED BY DIFFERENT ALPHA AND BETA CHAINS. BINDING BACTERIOCHLOROPHYLL MOLECULES, AND ARRANGED MOST PROBABLY IN TETRAMERIC STRUCTURES. DISCUSSED AROUND THE REACTION CENTER. THE N-TERMINAL GAMMA CHAINS MAY CONSTITUTE ADDITIONAL COMPONENTS."  
CC -1- STRECHLHAR LOCATION: Type II membrane protein. Inner membrane.  
CC PIP, S2164; S2164  
CC InterPro: IPR002361; Antenna.  
CC InterPro: IPR000666; Antenna\_a/b.  
CC Pfam: PF00566; LHC\_1  
CC F00566; F00566; ANTENNA\_COMP\_ALPHA; 1.  
CC Antenna complex; Light-harvesting polypeptide; Transmembrane;  
CC Magnesium; Bacteriochlorophyll; Inner membrane.  
CC CYTOPLASMIC (POTENTIAL).  
FT TRANSMEM 14 34  
FT TRANSMEW 14 34  
FT COMAIN 35 65  
FT METAL 29 29  
FT POTENTIAL.  
FT RESIDUALS (POTENTIAL).  
FT AXIAL LIGAND TO THE BACTERIOCHLOROPHYLL.  
FT MAGNESIUM (POTENTIAL).  
SQ SEQUENCE 65 AA: 7698 MW: 5544306/4443309A CRC64;  
Query Match 100.0%; Score 6; DB 1; Length 65;  
Best local similarity 50.0%; Pred No 78+02;  
Matches 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 WXXWKF 6  
DB 2 WXXWKF 7  
RESULT 2  
W99\_SEVIR

[illegible]

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05 entities requires a license agreement (see http://www.isb-sib.ch/announcement
06 or send an email to license@sib-sib.ch).
07 -----
08 EMBL accession: AAC07350.1
09 EMBL description: EMB07350.1, ALT_INT.
10 EMBL accession: EMB07350.1, ALT_INT.
11 EMBL accession: EMB07350.1, ALT_INT.
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99 EMBL accession: EMB07350.1, ALT_INT.
100 EMBL accession: EMB07350.1, ALT_INT.

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SEQUENCE R4 AA: 6811 MW: 21070P6267696B CP064;

Query Match 100.0% Score 6; FR 1; Length 84;

Best Local Similarity 50.0%; Pred. No. 9; SeqId2;

Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWVF 6

DB 63 WKHWF 68

RESULT 5

ACHA\_FELCA STANDARD; PRT; 84 AA.

ID ACHA\_FELCA STANDARD; PRT; 84 AA.

AC P54250; 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DT 15-JUN-2002 (Rel. 41, Last annotation update)

DR Acetylcholine receptor protein, alpha chain (Fragment)

GN CHRM1

OS Felis silvestris catus (Cat)

OC Eukaryota; Metazoa; Chordata; Mammalia; Carnivora; Felidae; Felis

OC Mammalia; Carnivora; Felidae; Felis

OC NCB1\_TaxID=9682

RP SEQUENCE FROM N A

RC TISSUE=Muscle;

RE MEDLINE=64147; PubMed=761917;

RT Baraban D, Oswald M, Fuchs S;

RT "The binding site of the nicotinic acetylcholine receptor in alpha1

RT Biochemistry 14:9172-9176(1995).

CC EXTENSION AFTER PIR IN ATTENTION. THE ACHR RECEPTOR IS AN

CC EXTENSIVE CHANGE IN COMPARISON THAT AFFECTS ALL SUBUNITS AND

CC LEADS TO OPENING OF AN ION-CONDUCTING CHANNEL ACROSS THE PLASMA

CC MEMBRANE

CC -1- SUBUNIT. FETTERER OF TWO ALPHA CHAINS, AND ONE EACH OF THE BETA,

CC DELTA, AND GAMMA (IN IMMATURE MUSCLE) OF EPHRIN (IN MATURE

CC MUSCLE) CHAINS.

CC -1- SUBUNIT. FETTERER OF TWO ALPHA CHAINS, AND ONE EACH OF THE BETA,

CC DELTA, AND GAMMA (IN IMMATURE MUSCLE) OF EPHRIN (IN MATURE

CC MUSCLE) CHAINS.

CC -1- SUBUNIT. FETTERER OF TWO ALPHA CHAINS, AND ONE EACH OF THE BETA,

CC DELTA, AND GAMMA (IN IMMATURE MUSCLE) OF EPHRIN (IN MATURE

CC MUSCLE) CHAINS.

CC -1- SUBUNIT. FETTERER OF TWO ALPHA CHAINS, AND ONE EACH OF THE BETA,

CC DELTA, AND GAMMA (IN IMMATURE MUSCLE) OF EPHRIN (IN MATURE

RESULT 6

YPI1\_ECOLI

AC P58095;

DT 16-OCT-2001 (Rel. 40, Created)

DT 15-JUN-2002 (Rel. 41, Last annotation update)

DR Hypothetical protein YPI1.

GN YPI1 OR E2641.1

OS Escherichia coli

OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;

OC Escherichia

OC NCB1\_TaxID=562;

RP SEQUENCE FROM N A

RC STRAIN=K12 / MG1655;

RE MEDLINE=942617; PubMed=9278503;

RT Blatter F, Plunkett G, III, Blitch C A, Perna N T, Burland V,

RT Piley M, Collado-Vides J, Glasner J D, Podge C V, Mayhew G F,

RT Granger T, Davis N W, Kirkpatrick H A, Goeden M A, Pusey D J,

RT Mau B, Shao Y;

RT "The complete genome sequence of Escherichia coli K-12"

RT Science 277:1233-1238(1997).

CC IDENTIFICATION, AND CONCEPTUAL TRANSLATION.

CC UNPUBLISHED OBSERVATIONS (JAN-1999).

CC SIMILARITY: STRONG TO E. coli YPI1

CC CAUTION: THIS IS A CONCEPTUAL TRANSLATION. A FRAME-SHIFT HAD TO BE

CC SUPPRESSED IN POSITIONS 8 AND 11.

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CC or send an email to [license@sib-sib.ch](mailto:license@sib-sib.ch).

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

CC EMBL, AE01345, , NCT\_A01345; CP064;

Query Match 100.0% Score 6; FR 1; Length 84;

Best Local Similarity 50.0%; Pred. No. 10; SeqId2;

Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWVF 6

DB 61 WRWVF 66

RESULT 7

YGS5\_MYCPN

AC P75147;

DT 16-OCT-2001 (Rel. 40, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DR Hypothetical lipid protein MEN650 precursor (E09\_01101)

GN MEN650 OR YP192

OS Mycoplasma pneumoniae

OC Bacteria; Firmicutes; Mollicutes; Mycoplasmatraceae; Mycoplasma

OC NCB1\_TaxID=2184;

RP SEQUENCE FROM N A

RC STRAIN=ATCC 29342 / M129;

RE MEDLINE=9710585; PubMed=8948633;

RT Himmelreich R, Hilbert H, Plogens H, Pirkl E, Li B.-C,

RT Hermann R;

RT "Complete sequence analysis of the genome of the bacterium Mycoplasma

RT





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CC or send an email to license@isb-sib.ch).
CC -----
CC DR EMBL: U32822; AAC3085.1; -.
CC TRIGR: H11436; -.
CC HYPERMUTATION: F111111; Cytidine deaminase.
CC SEQUENCE: 106 AA; 12273 MW; 405349980c4004c0064.
CC -----
Query Match
Best Local Similarity 50.0%; Score 6, DR 1, Length 100;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
OY 1 WXXWXXF 6
DB 48 WLSWVF 53

```

RESULT 10  
NIGM BOVIN  
ID NIGM BOVIN STANDARD. PPT: 109 AA  
AC 002374;  
DT 01-JUL-1993 (Rel. 26, Created)  
DT 01-JUL-1993 (Rel. 26, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE NADH-ubiquinone oxidoreductase ACQD subunit, mitochondrial precursor  
DE (EC 1.6.5.3) (P11644) (Complex I ACQD) (C1 ACQD3)  
GN NCBF82.  
OS Bos taurus (Bovine).  
OC Fungi; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eumetazoa; Cephalochordata; Primates; Rodentia, Muridae,  
OC Bovidae; Bovinae; Bos.  
RN NCB1\_TaxID 2911,  
RN [1]  
RP SEQUENCE FROM N. A. AND SEQUENCE OF 37 CDS.  
RT Tissue: Heart;  
RX MEDLINE: 9239317, PubMed: 1518244,  
RA Walker J.E., Arizumi T.M., Dupuis A., Fearney J.M., Fine M.,  
RA Med S.M., Pillington S.J., Panslow M.J., Skehel J.M.;  
FT "Sequences of 20 subunits of NADH-ubiquinone oxidoreductase from  
RT bovine heart mitochondria. Application of a novel strategy for  
sequencing proteins using the polymerase chain reaction."  
RL J. Mol. Biol. 276 (1993) 107-120;  
CC FUNCTION: TRANSFER OF ELECTRONS FROM NADH TO THE PEROXYRAC  
CHAIN. THE IMMEDIATE ELECTRON ACCEPTOR FOR THE FLUORE IS BELIEVED  
TO BE UBIQUINONE.  
CC -1- CATALYTIC ACTIVITY: NADH + ubiquinone = NAD(+) + ubiquinol  
CC -1- SUBUNIT: COMPLEX I IS COMPOSED OF ABOUT 40 DIFFERENT SUBUNITS.  
CC -1- SUBCELLULAR LOCATION: Mitochondrial inner membrane; matrix side.  
CC -----  
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CC -----
CC DR EMBL: X63216; CAA4901.1; -.
CC F1R: S28241; S28241.  
CC OVERLAPPING: NADH-ubiquinone, Mitochondria, Tissue: F111111.  
CC TRANSIT 1 36 MITOCHONDRIUM  
CC CHAIN 17 109 NADH-ubiquinone oxidoreductase ACQD  
CC SUBUNIT [1]  
CC FEATURE 106 AA; 12273 MW; 405349980c4004c0064.

```

DB 71 WLSWVF 76

```

RESULT 11  
ID HAVS CARAU STANDARD. PPT: 116 AA  
AC F19181;  
DT 01-NOV-1990 (Rel. 16, Created)  
DT 01-NOV-1990 (Rel. 16, Last sequence update)  
DT 15-JUL-1993 (Rel. 38, Last annotation update)  
DE Ig heavy chain V region 5A precursor.  
OS Carassius auratus (Goldfish).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Cyprinodontiformes;  
OC Cyprinidae; Carassius.  
OX NCB1\_TaxID=7957;  
RN NCB1 [1]  
RP SEQUENCE FROM N. A.  
RX MEDLINE=88144476; PubMed=3125551;  
RA Wilson M.R., Middleton D., Watt G.W.;  
FT "Amino acid sequence of heavy chain variable region 5A gene by direct structure  
RT and family relationships of two genes and a pseudogene in a teleost  
RT fish."  
PL Proc Natl Acad Sci U S A 86:1566-1570(1989).  
PF PIP: P29666; P29666.  
DE HSSP: P01722; 2F84.  
DE Interfer: IF803596, IG.MHC.  
DE Interfer: IF803596; IG.V.  
DR Pfam: PF00047; 1g.1.  
DR SMART, SMC310C, 1GV, 1.  
KW Immunoglobulin V region, Signal.  
FT SIGNAL 1 19  
FT CHAIN 20 116 IG HEAVY CHAIN V REGION 5A.  
FT DOMAIN 20 49 FRAMEWORK 1.  
FT DOMAIN 50 54 COMPLEMENTARITY-DETERMINING 1.  
FT DOMAIN 55 68 FRAMEWORK 2.  
FT DOMAIN 63 84 COMPLEMENTARITY-DETERMINING 2.  
FT DOMAIN 85 116 FRAMEWORK 3.  
FT DISUPTD 41 114 BY SIMILARITY.  
FT NCITER 116 116  
FT SEQUENCE 116 AA; 12908 MW; 922219822P199212 CIRC4;  
Query Match  
Best Local Similarity 50.0%; Score 6, DR 1, Length 100;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
OY 1 WXXWXXF 6  
DB 4 WLSWVF 9

RESULT 12  
ID MERT STAU  
ID MERT STAU STANDARD. PPT: 128 AA.  
AC P08656;  
DT 01-JAN-1988 (Rel. 06, Created)  
DT 01-JAN-1988 (Rel. 06, Last sequence update)  
DT 01-NOV-1997 (Rel. 35, Last annotation update)  
DE Mercury transport protein (Mercury ion transport protein).  
GN MERT.  
OS Staphylococcus aureus.  
OC Bacteria; Firmicutes; Bacillales; Staphylococcus.  
CC F1458.  
CC NCB1\_TaxID=1280;  
RN NCB1 [1]  
RP SEQUENCE FROM N. A.  
RX MEDLINE=87260937; PubMed=3037534;  
PA Laddaga P.A., Chu L., Mista T.V., Silver S.;  
FT "Molecular sequence and expression of the mercurial protein gene  
RT operon from Staphylococcus aureus plasmid p128."  
RL Proc Natl Acad Sci U S A 84:5106-5110(1987).  
CC FUNCTION: INVOLVED IN MERCURY TRANSPORT. PASSES A HG(II) ION  
CC FROM THE PERIPLASMIC MERT PROTEIN TO THE MEMBRANE PROTEIN.



```

ID  YA91 MYCPN      STANDARD;      PRT;      138 AA.
AC  P75602;
DT  16-OCT-2001 (Rel. 40, Created)
DT  16-OCT-2001 (Rel. 40, Last sequence update)
DT  16-OCT-2001 (Rel. 40, Last annotation update)
DE  Hypothetical protein MPN091 (R02_orf138).
GN  MPN091 ORF MP064.
OS  Mycoplasma pneumoniae.
OC  Bacteria; Firmicutes; Mollicutes; Mycoplasmataceae; Mycoplasma.
OX  NCBI_TaxID=2104;
RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN-ATCC 29342 / M129;
RX  MEDLINE:97105885; PubMed:8948633;
RA  Hammelreich R., Hilbert H., Pliagens H., Pirki E., Li B.-C.,
RA  Herrmann R.;
RT  "Complete sequence analysis of the genome of the bacterium Mycoplasma
RT  pneumoniae."
RL  Nucleic Acids Res. 24:4420-4449(1996).
CC  - SIMILARITY: ALMOST IDENTICAL TO M.FRECMONTIAE MEN413 AND MEN463.
CC  -----
CC  This SWISS-PROT entry is copyright. It is produced through a collaboration
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CC  or send an email to license@sib-sib.ch).
CC  -----
CC  EMBL; AE000008; AAB95711.1; -
CC  KW  Hypothetical protein; Complete proteome;
CC  SEQUENCE 138 AA, 15723 MW, F52380C428F43709 DCD64;
SQ

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```

Query Match          100.0%; Score 6; DB 1; Length 138;
Best Local Similarity 50.0%; Pval No. 1.4e+03;
Matches 3; Conservation 3; Mismatches 0; Indels 0; Gaps 0;

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QY  1 WXXXXF 6
    |||||
DB  36 MSGMSF 41

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Search completed: February 20, 2003, 11:24:57  
 Job time : 12.3333 secs





[illegible]

DT 01-OCT-2000 (TREMURel 15, Created)  
 DT 01-OCT-2000 (TREMURel 15, Last sequence update)  
 DT 01-DEC-2001 (TREMURel 19, Last annotation update)  
 DE FAMAM1 (Fragment)  
 GN FAMAM1  
 OS Homo sapiens (Human)  
 CC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo;  
 CC NRI TaxID=9606;  
 RN [1] TaxID=9606;  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE-20349292; PubMed10989047;  
 RA Vincent J.B., Hepburn J.A., Gurling H.M.D., Bolton P.F., Roberts W.,  
 RA Scherer S.W.;  
 RT Identification of a novel gene on chromosome 7q31 that is interrupted  
 RT by a translocation breakpoint in an autistic individual.;  
 RL Am. J. Hum. Genet. 67:510-514(2001).  
 DR EMBL: AF234886; AAF8549.1; -;  
 FT NON-TER 50  
 FT 50  
 SQ SEQUENCE 50 AA, 1041 MW, 16490.724999999999 C7664,  
 Query Match, 100.0%, Score 6, DR 4, Length 50,  
 Best Local Similarity 50.0%, Pred. No. 2, Total,  
 Matches 3, Conservative 3, Mismatches 0, Indels 0, Gaps 0,  
 QY 1 WXXXXF 6  
 Db 24 WYWWF 29  
 RESULT 7  
 Q9PH20 PRELIMINARY: PRT, 59 AA.  
 ID Q9PH20  
 AC Q9PH20  
 DT 01-OCT-2000 (TREMURel 15, Created)  
 DT 01-OCT-2000 (TREMURel 15, Last sequence update)  
 DT 01-JUN-2002 (TREMURel 21, Last annotation update)  
 DE HYPOHETICAL PROTEIN XE0126.  
 GN XE0126.  
 OS Xylella fastidiosa.  
 CC Bacteria; Proteobacteria; gamma subdivision; Xanthomonas group;  
 CC Xylella.  
 CC Xylella fastidiosa.  
 RN [1] TaxID=2071.  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE-20365717; PubMed16918347;  
 RA Simpson A.J.G., Peinath F.C., Atirah F., Atreya F.A., Ascarin M.,  
 RA Alvarado R., Alves L.M., Ataya J.F., Balag S., Papista C.S.,  
 RA Barrios M.H., Bonaccorsi A.D., Bordin S., Bova J.M., Britton M.P.S.,  
 RA Bueno M.P., Camargo A.A., Camargo L.E.A., Carraro D.M., Carraro H.,  
 RA Colares R.R., Colares C., Costa F.F., Costa M.G.P., Costa-Mello C.M.,  
 RA Coutinho L.L., Cristofani W., Dias-Nova E., Dorschner G.E., Dory H.,  
 RA Facinanci A.P., Ferreira A.D.S., Ferreira V.C.A., Ferra D.,  
 RA Fraga J.S., Frasca S.G., Franco M.C., Frohme M., Furian L.P.,  
 RA Gattner R., Goldman G.H., Goldman M.H.S., Gomes S.L., Graber A.,  
 RA Ho P.Y., Hohenstein J.D., Hohenstein M.E., Kasper E.L., Kijak J.P.,  
 RA Kleber J.F., Krumm F.E., Laigret F., Lamballe M.P., Leite L.C.G.,  
 RA Lemos M.G.M., Lemos M.V.F., Lopes M.A., Lopes C.P., Machado J.A.,  
 RA Machado M.A., Madeira A.M.B., Madeira H.M.F., Marino C.I.,  
 RA Marques M.V., Martins E.A.L., Martins E.M.F., Matoskova A.V.,  
 RA Marwick C.F.M., Miranda F.C., Miyaki C.V., Monteiro-Vitorino C.R.,  
 RA Moon D.H., Nagai M.A., Nascimento A.L.T.C., Netto L.P.S.,  
 RA Nishi A.C.J., Nobrega F.G., Nunes L.F., Oliveira M.A.,  
 RA de Oliveira M.C., de Oliveira P.C., Pulieri P.A., Reis A.,  
 RA Peixoto P.F., Pereira G.A.G., Pereira H.A.Jr., Pequeno J.B.,  
 RA Quintao P.R., Roberto P.G., Rodrigues V., de Rosa A.U.M.,  
 RA de Rosa V.B.Jr., de Sa F.G., Sanzini F.V., Sawasji H.E.,  
 RA da Silva A.C.R., da Silva A.M., da Silva F.R., Silva W.A.Jr.,  
 RA da Silveira C.F., Silvestri M.L., Silveira M.J., de Souza A.A.,  
 RA de Souza A.P., Terenzi M.F., Truffi D., Tsai S.M., Tshako M.H.,  
 RA Valodia H., Van Sluys M.A., Verjovsky-Almeida S., Vitorino A.L.,  
 RA Zago M.A., Zatz M., Zetianis J., Zetianis J.C.,

RT "The genome sequence of the plant pathogen *Xylella fastidiosa*."  
 RL Nature 406:151-159(2000).  
 DR EMBL: AF003865; AAF8293.1; -;  
 DR InterPro: IPR001230; Pfam: site.  
 DR PROSITE: PS00294; PENNYLATION; GMPNMN.1.  
 KW Hypothetical protein; Complete proteome.  
 SQ SEQUENCE 53 AA, 7368 MW, 12704.425000000000 C7664;  
 Query Match, 100.0%, Score 6, DR 15, Length 59,  
 Best Local Similarity 50.0%, Pred. No. 3, Total,  
 Matches 3, Conservative 3, Mismatches 0, Indels 0, Gaps 0,  
 QY 1 WXXXXF 6  
 Db 54 WRCWLF 59  
 RESULT 8  
 ID Q95619 PRELIMINARY: PRT, 61 AA.  
 AC Q95619;  
 DT 01-NOV-1998 (TREMURel 09, Created)  
 DT 01-NOV-1998 (TREMURel 09, Last sequence update)  
 DT 01-MAR-2002 (TREMURel 20, Last annotation update)  
 DE L0012 (Hypothetical 5.5 kDa protein).  
 GN 25095 OR ECS4544.  
 OS Escherichia coli, and  
 OS Escherichia coli O157:H7.  
 CC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;  
 CC Escherichia.  
 CC NCB1 TaxID=562, 83334;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX SPECIES-E COLI: STRAIN-EDL933;  
 RX MEDLINE-20320985; PubMed9673266;  
 RA Perna N.T., Mayhew G.F., Postal G., Elliott G., DeLong-Eby M.S.,  
 RA Perna J.B., Mayhew G.F., Postal G., Elliott G., DeLong-Eby M.S.,  
 RA Kaper J.B., Blatterer F.R.,  
 RT "Molecular evolution of a pathogenicity island from enterohemorrhagic  
 RT Escherichia coli O157:H7.";  
 RL Infect. Immun. 66:3810-3817(1998).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RX SPECIES-E COLI: STRAIN-O157:H7 / EDL933 / ATCC 700677;  
 RX MEDLINE-21074935; PubMed11206551;  
 RA Perna N.T., Plunkett G. III, Burland V., Mau B., Glasner J.D.,  
 RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,  
 RA Paulsen O., Hackett J., Klink S., Boulton A., Shao Y., Miller L.,  
 RA Drotbock A.C., Davis N.M., Lin A., Leland E.T., Peterson K.,  
 RA Apolack J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,  
 RA Welch P.A., Blatterer F.R.,  
 RT "Genome sequence of enterohemorrhagic Escherichia coli O157:H7.";  
 RL Nature 409:529-533(2001).  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RX SPECIES-E COLI: STRAIN-O157:H7 / P1W 0509950;  
 RX MEDLINE-2156031; PubMed11258796;  
 RA Hayashi T., Makino K., Chitishi Y., Yokoyama Y.,  
 RA Hara G., Chitishi E., Nakayama Y., Morita T., Tanaka M., Tate T.,  
 RA Tada T., Takai H., Honda T., Sasaki K., Ogatawara N., Yasunaga T.,  
 RA Kohara S., Shibata T., Harashina M., Shitogawa H.,  
 PT Complete genome sequence of enterohemorrhagic Escherichia coli  
 PT O157:H7 and genomic comparison with a laboratory strain O157.  
 PT DNA Res. 8:11-22(2001).  
 DR EMBL: AF071034; AAC349.1; -;  
 DR EMBL: AF071034; AAC349.1; ALT-INT  
 DR EMBL: AF002566; BAB37967.1; -;  
 KW Hypothetical protein; Complete proteome.  
 SQ SEQUENCE 51 AA, 7286 MW, 59748.116500000000 C7664;  
 Query Match, 100.0%, Score 7, DR 16, Length 61,  
 Best Local Similarity 50.0%, Pred. No. 3, Total,  
 Matches 3, Conservative 3, Mismatches 0, Indels 0, Gaps 0,

QY 1 WXXXWF 6  
DB 3 WDWRAF 8

## RESULT 9

Q9HWK5 PRELIMINARY, FRT, 64 AA.

AC Q9HWK5  
DT 01-MAR-2003 (TREMBL: 20, last sequence update)  
DI 01-MAR-2003 (TREMBL: 20, last sequence update)  
DE Hypothetical protein (Fragment)  
OS Homo sapiens (Human)  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eulalia; Placental; Catarrhini; Hominoidea; Homo;  
NCBI\_TaxId:9606;  
RN 1  
RP SEQUENCE FROM N.A.  
RC TISSUE-BRAIN;  
RA Submitted (DEC-2001) to the EMBL/GenBank/DDBJ databases.  
DR EMBL: BC017733; ANH17733.1;  
KW Hypothetical protein  
FT NON\_TER  
SQ SEQUENCE 64 AA, 7480 MW, 117582ATTAGGCTG GCTG,  
Best Local Similarity 50.0%; Score 6; DB 4; Length 64;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXWF 6  
DB 16 WDWRAF 21

## RESULT 10

Q9EXE4 PRELIMINARY, FRT, 70 AA.

AC Q9EXE4  
DT 01-MAR-2003 (TREMBL: 16, last sequence update)  
DI 01-MAR-2003 (TREMBL: 16, last sequence update)  
DE Putative thymidylate synthase (Fragment)  
OS Bacillus subtilis  
OC Bacteria; Firmicutes; Bacilli; Clostridia; Bacillales; Bacillaceae; Bacillus  
NCBI\_TaxId:1284;  
RN 1  
RP SEQUENCE FROM N.A.  
RC Strain 15A21;  
RA Basso A.; Campanile G.; Ferrara L.; Damiani G.;  
RT "Sequencing and analysis of DNA fragments obtained by primer walking  
polymorphic DNA (PCR) technique from *Bacillus subtilis* strain 15A21;  
J. Clin. Microbiol. 40:1027-1030 (2002)."  
DR EMBL: AU046222; CMO20646.1;  
DR HSSP: P03469; TSY;  
DR InterPro: IPR000100; Thymidylt SYN  
DR Pfam: PF00303; Thymidylat SYN\_1  
DR ProDom: PD001190; Thymidylt SYN\_1  
FT NON\_TER  
SQ SEQUENCE 70 AA, 5183 MW, 42019 GCTTCTG GCTG,  
Best Local Similarity 50.0%; Score 6; DB 4; Length 70;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXWF 6  
DB 60 WDWRAF 65

## RESULT 11

Q9HST8 PRELIMINARY, FRT, 71 AA.

AC Q9HST8  
DT 01-MAR-2003 (TREMBL: 16, last sequence update)  
DI 01-MAR-2003 (TREMBL: 16, last sequence update)  
DE Hypothetical protein (Fragment)  
OS Homo sapiens (Human)  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eulalia; Placental; Catarrhini; Hominoidea; Homo;  
NCBI\_TaxId:9606;  
RN 1  
RP SEQUENCE FROM N.A.  
RC KEDINE2050493; PubMed:1016950;  
RA KJ W.Y.; Kennedy S.P.; Mahalis G.G.; Bergquist B.; Fan M.;  
RA Shalla H.D.; Lasky G.R.; Ealla N.S.; Thompson V.; Shogna J.;  
RA Swartzell S.; Weir D.; Hall G.; Hall T.A.; Weir D.; Goo Y.A.;  
RA Lehnus F.; Folger F.; Tyr P.; Hansen M.; Hough D.W.;  
RA Wallick D.G.; Altshuler E.; Krebs M.F.; Argenteau C.M.; Dale H.;  
RA Isenberger T.A.; Peck R.F.; Polischud M.; Spudis O.B.; Song R.H.;  
RA Alam M.; Fieles T.; Holt S.; Daniels C.D.; Dennis P.P.; Meyer A.D.;  
RA Ehardt H.; Lise T.M.; Ling F.; Ploeg Y.; Wood L.; Dacarta C.;  
FT Complete sequence of *Halobacterium salinarum* R1.  
RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181 (2000).  
KW Complete proteome.  
SQ SEQUENCE 71 AA, 8002 MW, 157341TTCGATG CAGG,  
Best Local Similarity 50.0%; Score 6; DB 17; Length 71;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXWF 6  
DB 49 WDWRAF 54

## RESULT 12

Q50168 PRELIMINARY, FRT, 74 AA.

AC Q50168  
DT 01 NOV 1996 (TREMBL: 01, last sequence update)  
DI 01 NOV 1996 (TREMBL: 01, last sequence update)  
DE Mycobacterium leprae  
OC Bacteria; Firmicutes; Actinobacterii; Actinobacteriales;  
OC Actinomycetales; Corynebacteriiformes; Mycobacteriales; Mycobacterium;  
OS NCBI\_TaxId:1769;  
RN 1  
RP SEQUENCE FROM N.A.  
RC Smith D.R.;  
RT "Isolation and characterization of *Mycobacterium leprae* strains;  
J. Clin. Microbiol. 34:1027-1030 (1996)."  
DR EMBL: U00097; AA06314.1;  
DR EMBL: U00097; AA06314.1;  
SQ SEQUENCE 74 AA, 5043 MW, 32000A027TCTG TCGG,  
Best Local Similarity 50.0%; Score 6; DB 4; Length 74;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXWF 6  
DB 59 WDWRAF 64

## RESULT 13



```

058793
ID 068793 PRELIMINARY; PRT: 76 AA.
AC 068793;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-APR-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-MAR-2002 (TrEMBLrel. 20, Last annotation update)
DE Hypothetical protein Y1001
GN Y1001 OR YPMT1.60C.
OS Versinia pestis.
OC Plasmid pMT1 (pMT-1).
OC Paerctria, Proteobacteria, gamma subdivision, Enterobacteriaceae,
OC Versinia.
OX NCBI TaxID=632;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-KIM5;
RX MEDLINE:98422474; PubMed=9748454;
RA Hu P., Elliott J., McCreedy P., Skowronski E., Ganes J.,
RA Kobayashi A., Brubaker R.R., Garcia E.;
RT "Structural organization of virulence-associated plasmids of Versinia
RT pestis";
RL J. Bacteriol. 190:5192-5202 (1998)
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-KIM10;
RX MEDLINE:9943898; PubMed=9926348;
RA Lindler I.R., Plano G.V., Burland V., Mayhew G.F., Plattner F.P.;
RT "Complete DNA sequence and detailed analysis of the Versinia pestis
RT KIM5 plasmid encoding murine toxin and capsular antigen";
RL Infect. Immun. 66:5741-5742 (1998)
RN [3]
RP SEQUENCE FROM N.A.
PT STRAIN-0042 / HIVAR ORIENTALIS; PLASMID-FMT1 (pMT 1);
RX MEDLINE:21470413; PubMed=11586360;
RA Parkhill J., Wren B.W., Thomson N.P., Tishall P.W., Halsey N.T.G.,
RA Penrice M.B., Sebatina M., James P.D., Churruarin C., Whittall P.T.,
RA Baker S., Basham P., Bentley S.P., Brooks K., Cardeno-Tarrega A.M.,
RA Chillingworth T., Cronin A., Davies P.M., Davis F., Dougan G.,
RA Fellwell T., Hamlin N., Helgryd S., James K., Karyshev A.V.,
RA Leather S., Moulé S., Oyston P.C.F., Quail M., Rutherford K.,
RA Simmonds M., Skelton J., Stevens K., Whitehead S., Barrell P.G.;
RT "Genome sequence of Versinia pestis: the causative agent of plague";
RL Nature 413:523-527 (2001);
DR EMBL: AF051947; AAC13243.1;
DR EMBL: AF074611; AAC82749.1;
DR EMBL: AL117211; CAB55242.1;
KW Hypothetical protein; Plasmid; Complete proteome.
SQ SEQUENCE 76 AA; 8807 MW; 7EEF4937755036EB CRC64;

Query Match 100.0%; Score 6; DB 16; Length 76;
Best Local Similarity 50.0%; Pred. No. 3, 9e+03;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 50 WRTWNP 55

RESULT 14
09S4W3
ID 09S4W3 PRELIMINARY; PRT: 77 AA.
AC 09S4W3;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-MAY-2000 (TrEMBLrel. 13, Last annotation update)
DE Hypothetical 9.1 kDa protein.
OS Escherichia coli.
OC Plasmid F.
OC Paerctria, Proteobacteria, gamma subdivision, Enterobacteriaceae,
OC Escherichia.
OX NCBI TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.

```

```

EX MEDLINE:99296678; PubMed 10365527;
RA Manwaring N.P., Skurray P.A., Fitch N.;
RT "Nucleotide sequence of the F plasmid leading region";
RL Plasmid 41:219-225 (1999);
DR EMBL: AF105323; AAD47188.1;
KW Hypothetical protein; Plasmid.
SQ SEQUENCE 77 AA; 9083 MW; 2645F03579477A CRC64;

Query Match 100.0%; Score 6; DB 2; Length 77;
Best Local Similarity 50.0%; Pred. No. 4e+03;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 53 WRTWNP 58

RESULT 15
09WTF7
ID 09WTF7 PRELIMINARY; PRT: 79 AA.
AC 09WTF7;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE YdeA protein.
GN YDEA.
OS Escherichia coli.
OC Plasmid p100.
OC Paerctria, Proteobacteria, gamma subdivision, Enterobacteriaceae,
OC Escherichia.
OX NCBI TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.
RA Sampei G., Mizobuchi K.;
RT "Organization and classification of plasmid genomes: complete
RT nucleotide sequence of the p100 genome";
PL Submitted (MAY-1999) to the EMBL/Genbank/DBJ databases.
DR EMBL: AF000140; PA079837.1;
KW SEQUENCE 79 AA; 9368 MW; 7107949777777777 CRC64;

Query Match 100.0%; Score 6; DB 3; Length 79;
Best Local Similarity 50.0%; Pred. No. 4e+03;
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXXXF 6
DB 55 WRTWNP 60

Search completed: February 20, 2003, 11:28:57
Job time: 47 secs

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GenCore version 5.1.3  
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OW protein - protein search, using fw model

Run on: February 20, 2003, 11:31:07, Search time 29.3333 seconds  
(without alignments)  
27,256 Million cell updates/sec

Title: US-09-912-414-9

Perfect score: 6

Sequence: 1 WXXWXP

Sorting table: GILPCGX

Gap: 60.0, Gapext: 60.0

Searched: 908470 seqs, 13350620 residues

Word size: 9

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum hit seq length: 200000000

Post-processing: Listing first 45 summaries

Database:

1: /SID82/gcgdata/geneseq/geneexp-emb1/AA1980.DAT:\*  
2: /SID82/gcgdata/geneseq/geneexp-emb1/AA1981.DAT:\*  
3: /SID82/gcgdata/geneseq/geneexp-emb1/AA1982.DAT:\*  
4: /SID82/gcgdata/geneseq/geneexp-emb1/AA1983.DAT:\*  
5: /SID82/gcgdata/geneseq/geneexp-emb1/AA1984.DAT:\*  
6: /SID82/gcgdata/geneseq/geneexp-emb1/AA1985.DAT:\*  
7: /SID82/gcgdata/geneseq/geneexp-emb1/AA1986.DAT:\*  
8: /SID82/gcgdata/geneseq/geneexp-emb1/AA1987.DAT:\*  
9: /SID82/gcgdata/geneseq/geneexp-emb1/AA1988.DAT:\*  
10: /SID82/gcgdata/geneseq/geneexp-emb1/AA1989.DAT:\*  
11: /SID82/gcgdata/geneseq/geneexp-emb1/AA1990.DAT:\*  
12: /SID82/gcgdata/geneseq/geneexp-emb1/AA1991.DAT:\*  
13: /SID82/gcgdata/geneseq/geneexp-emb1/AA1992.DAT:\*  
14: /SID82/gcgdata/geneseq/geneexp-emb1/AA1993.DAT:\*  
15: /SID82/gcgdata/geneseq/geneexp-emb1/AA1994.DAT:\*  
16: /SID82/gcgdata/geneseq/geneexp-emb1/AA1995.DAT:\*  
17: /SID82/gcgdata/geneseq/geneexp-emb1/AA1996.DAT:\*  
18: /SID82/gcgdata/geneseq/geneexp-emb1/AA1997.DAT:\*  
19: /SID82/gcgdata/geneseq/geneexp-emb1/AA1998.DAT:\*  
20: /SID82/gcgdata/geneseq/geneexp-emb1/AA1999.DAT:\*  
21: /SID82/gcgdata/geneseq/geneexp-emb1/AA2000.DAT:\*  
22: /SID82/gcgdata/geneseq/geneexp-emb1/AA2001.DAT:\*  
23: /SID82/gcgdata/geneseq/geneexp-emb1/AA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	6	100.0	6	15	AA857386
2	6	100.0	6	15	AA857391
3	6	100.0	6	21	AA801492
4	6	100.0	6	21	AA801497
5	6	100.0	6	21	AA801499
6	6	100.0	6	21	AA801505
7	6	100.0	6	21	AA801506
8	6	100.0	6	21	AA801508
9	6	100.0	7	22	AAE12181
10	6	100.0	7	22	AA845777

11	6	100.0	8	15	AA857399	Peptide for treati
12	6	100.0	9	21	AA801498	Peptide which bind
13	6	100.0	9	21	AA801499	Peptide for treati
14	6	100.0	10	15	AA857400	Peptide for treati
15	6	100.0	11	19	AA862149	Peptide for treati
16	6	100.0	11	21	AA801505	Peptide which bind
17	6	100.0	12	15	AA857401	Peptide for treati
18	6	100.0	12	21	AA801504	Peptide which bind
19	6	100.0	13	21	AA801505	Peptide which bind
20	6	100.0	13	18	AA801506	Peptide which bind
21	6	100.0	13	18	AA801507	Peptide which bind
22	6	100.0	14	22	AA801508	Peptide which bind
23	6	100.0	15	20	AA801509	Peptide which bind
24	6	100.0	15	21	AA801510	Peptide which bind
25	6	100.0	15	21	AA801511	Peptide which bind
26	6	100.0	15	21	AA801512	Peptide which bind
27	6	100.0	15	21	AA801513	Peptide which bind
28	6	100.0	17	21	AA801514	Peptide which bind
29	6	100.0	19	22	AA801515	Peptide which bind
30	6	100.0	20	23	AA801516	Peptide which bind
31	6	100.0	20	23	AA801517	Peptide which bind
32	6	100.0	20	23	AA801518	Peptide which bind
33	6	100.0	20	23	AA801519	Peptide which bind
34	6	100.0	21	22	AA801520	Peptide which bind
35	6	100.0	22	22	AA801521	Peptide which bind
36	6	100.0	22	22	AA801522	Peptide which bind
37	6	100.0	22	22	AA801523	Peptide which bind
38	6	100.0	22	22	AA801524	Peptide which bind
39	6	100.0	22	22	AA801525	Peptide which bind
40	6	100.0	22	22	AA801526	Peptide which bind
41	6	100.0	22	22	AA801527	Peptide which bind
42	6	100.0	22	22	AA801528	Peptide which bind
43	6	100.0	22	22	AA801529	Peptide which bind
44	6	100.0	22	22	AA801530	Peptide which bind
45	6	100.0	22	22	AA801531	Peptide which bind

#### ALIGNMENTS

RESULT 1  
ID AA857386 standard, peptide: 6 AA.  
XX AA857386;  
AC  
DT 21-MAR-1995 (first entry)  
XX  
DE Peptide for treating diseases related to anti DNA antibodies.  
XX  
KW Carrier, adsorbing agent, treatment, anti-DNA antibody, immunological.  
XX  
OS Synthetic  
XX  
PR JF06192290-A.  
XX  
PD 12-JUL-1994.  
XX  
PF 18-JAN-1993; 93JP-0006098.  
XX  
FR 30-SEP-1992; 93JP-0261821.  
XX  
PA (KIPS) KIPAPAY CO LTD  
XX  
DR WPI, 1994-260510/32.  
XX  
PT A peptide and an adsorbing agent prepd. by immobilising it on a  
PT carrier - useful for treatment of diseases related to anti DNA  
PT antibodies and immune complexes  
XX  
PS Disclosure: Page 11; 14pp; Japanese.  
XX  
CC The sequences given in AA857386-413 are peptides which are all covered

CC by the claimed generic formula.  
 CC H X (A,B) Y Z  
 CC A = Trp, Phe or a peptide fragment consisting of 2 residues;  
 CC B = Trp, Phe, Asn or Glu;  
 CC X and Y = a bond or Asp, Glu, Arg, Lys, His or a peptide fragment  
 CC consisting of 2-10 residues, provided that at least one of  
 CC X or Y are present;  
 CC Z = OH or NH<sub>2</sub>; and  
 CC n = 2-5.  
 CC These peptides may be immobilised on a carrier in the preparation of an  
 CC absorbing agent which may be used in the treatment of diseases related  
 CC to anti-DNA antibodies and/or immune complex.  
 CC Sequence 6 AA;  
 CC  
 CC Query Match 100.0%; Score 6; DB 15; Length 6;  
 CC Best local similarity 100.0%; Identical 100.0%;  
 CC Mismatch 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC  
 CC QY 1 WXXWXP 6  
 CC 111111  
 CC 1 WWRWMP 6  
 CC  
 CC RESULT 2  
 CC AAB57391  
 CC ID AAB57391 standard; Protein; 6 AA.  
 CC  
 CC AC AAB57391;  
 CC  
 CC DT 21-MAP-1998 (first entry)  
 CC  
 CC XX Peptide for treating diseases related to anti-DNA antibodies.  
 CC DE  
 CC XX Carrier, absorbing agent, treatment, anti-DNA antibody, immune complex.  
 CC XX  
 CC OS Synthetic  
 CC  
 CC PR Cited in: A.  
 CC XX  
 CC PD 12-000-1994.  
 CC  
 CC PF 18-JAN-1993; 982P-0000098.  
 CC  
 CC PR 11-DEF-1993; 982P-0000098.  
 CC  
 CC PA (KORS) KIRAPAY 0017P  
 CC  
 CC XX WT1, 1994 06110101.  
 CC  
 CC XX A peptide and an absorbing agent, provided by immobilising it on a  
 CC carrier - useful for treatment of diseases related to anti-DNA  
 CC antibodies and immune complexes  
 CC  
 CC PS Disclosure; Page 11; 14pp; Japanese  
 CC  
 CC XX The sequences given in AAB57391-413 are preferred which are all covered  
 CC by the claimed generic formula:  
 CC H X (A,B) Y Z  
 CC A = Trp, Phe or a peptide fragment consisting of 2 residues;  
 CC B = Trp, Phe, Asn or Glu;  
 CC X and Y = a bond or Asp, Glu, Arg, Lys, His or a peptide fragment  
 CC consisting of 2-10 residues, provided that at least one of  
 CC X or Y are present;  
 CC Z = OH or NH<sub>2</sub>; and  
 CC n = 2-5.  
 CC These peptides may be immobilised on a carrier in the preparation of an  
 CC absorbing agent which may be used in the treatment of diseases related  
 CC to anti-DNA antibodies and/or immune complex.  
 CC Sequence 6 AA;  
 CC  
 CC Query Match 100.0%; Score 6; DB 15; Length 6;

CC Best local similarity 100.0%; Identical 100.0%;  
 CC Mismatch 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC  
 CC QY 1 WXXWXP 6  
 CC 111111  
 CC 1 WWRWMP 6  
 CC  
 CC RESULT 3  
 CC AAB01492  
 CC ID AAB01492 standard; peptide; 6 AA.  
 CC  
 CC AC AAB01492;  
 CC  
 CC DT 08-NOV-2000 (first entry)  
 CC  
 CC XX For the treatment of diseases related to anti-DNA antibodies.  
 CC DE  
 CC XX DNA binding, transcription factor, E2F, E2F-1, cell cycle, DP-1,  
 CC activation, transcription, apoptosis, proliferation disorder,  
 CC psoriasis, restenosis.  
 CC  
 CC OS Synthetic.  
 CC  
 CC PN WO200044771-A1.  
 CC  
 CC PD 03-AUG-2000.  
 CC  
 CC PF 26-JAN-2000; 2000W-0800227.  
 CC  
 CC PR 26-JAN-1999; 982P-0000710.  
 CC  
 CC FA (FROD) FRODIFIX LTD.  
 CC  
 CC ST Mueller F, Koenigsmann RE, Montignani S;  
 CC  
 CC XX WPI, 2000 020006/48.  
 CC  
 CC XX Peptides binding to the DNA binding domain of transcription factor E2F  
 CC and inhibiting cell cycle progression, useful for the treatment of  
 CC cancer  
 CC  
 CC PS Claim 6, Page 2, 42pp, English.  
 CC  
 CC XX Peptides which bind to the DNA binding domain of transcription  
 CC factor E2F and inhibit cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC at the activation of transcription by E2F. They may also be used to  
 CC study the role of E2F in cell cycle arrest in  
 CC a cell, particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis.  
 CC  
 CC SQ Sequence 6 AA;  
 CC  
 CC Query Match 100.0%; Score 6; DB 21; Length 6;  
 CC Best local similarity 100.0%; Identical 100.0%;  
 CC Mismatch 0; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC  
 CC QY 1 WXXWXP 6  
 CC 111111  
 CC 1 WWRWMP 6  
 CC  
 CC RESULT 4  
 CC AAB01497  
 CC ID AAB01497 standard; peptide; 6 AA.  
 CC  
 CC AC AAB01497;  
 CC  
 CC DT 09-NOV-2000 (first entry)  
 CC  
 CC XX Peptide which binds to transcription factor E2F 1 DNA binding domain.  
 CC DE  
 CC XX

KW DNA binding, transcription factor; E2F, E2F-1; cell cycle; DP-1;  
 KW activation; transcription, apoptosis, proliferative disorder,  
 KW psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 FH Key Location/Qualifiers  
 FT Misc-difference 2 /note= "Any amino acid"  
 FT Misc-difference 3 /note= "Any amino acid"  
 FT Misc-difference 5 /note= "Any amino acid"  
 FT Misc-difference 5 /note= "Any amino acid"  
 PN WO200044771-A1.  
 PD 03-AUG-2000.  
 XX  
 XX 26-JAN-2000; 2000WO-GB00227.  
 PR 26-JAN-1999; 99GB-0001710.  
 XX  
 XX (PROL-) PROLIFIX LTD.  
 PI Mueller R, Kontermann RE, Montigiani S;  
 XX  
 XX WPI; 2000-532806/48.  
 DR Peptides binding to the DNA binding domain of transcription factor E2F  
 PT and inhibiting cell cycle progression, useful for the treatment of  
 PT cancer  
 PT  
 PS Claim 4; Page 9; 42pp; English.  
 XX  
 CC Peptides which bind to the DNA binding domain of transcription  
 CC factor E2F and inhibit cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC or the activation of transcription by E2F-1/DP-1 heterodimers. They  
 CC may also be used for inducing apoptosis and/or cell cycle arrest in  
 CC a cell, particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis.  
 CC  
 XX  
 SQ Sequence 6 AA;  
 Query Match 100.0%; Score 6; DB 21; Length 6;  
 Best Local Similarity 100.0%; Prod No 7 Re:05;  
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWXF 6  
 DB 1 WXXWXF 6  
 RESULT 5  
 AAB01499  
 ID AAB01499 standard; peptide; 6 AA.  
 XX  
 AC AAB01499;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DF Peptide which binds to transcription factor E2F-1 DNA binding domain.  
 XX  
 KW DNA binding; transcription factor; E2F, E2F-1; cell cycle; DP-1;  
 KW activation; transcription; apoptosis; proliferative disorder;  
 KW psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 FH Key Location/Qualifiers  
 FT Misc-difference 2 /note= "Any amino acid"  
 FT Misc-difference 3 /note= "Any amino acid"  
 FT Misc-difference 3 /note= "Any amino acid"

FT /note= "Any amino acid"  
 XX  
 XX WO200044771-A1.  
 XX  
 PD 03-AUG-2000.  
 XX  
 XX 26-JAN-2000; 2000WO-GB00227.  
 PR 26-JAN-1999; 99GB-0001710.  
 XX  
 XX (PROL-) PROLIFIX LTD.  
 PI Mueller R, Kontermann RE, Montigiani S;  
 XX  
 XX WPI; 2000-532806/48.  
 DR Peptides binding to the DNA binding domain of transcription factor E2F  
 PT and inhibiting cell cycle progression, useful for the treatment of  
 PT cancer  
 PT  
 PS Claim 4; Page 9; 42pp; English.  
 XX  
 CC Peptides which bind to the DNA binding domain of transcription  
 CC factor E2F and inhibit cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC or the activation of transcription by E2F-1/DP-1 heterodimers. They  
 CC may also be used for inducing apoptosis and/or cell cycle arrest in  
 CC a cell, particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis.  
 CC  
 XX  
 SQ Sequence 6 AA;  
 Query Match 100.0%; Score 6; DB 21; Length 6;  
 Best Local Similarity 81.4%; Prod No 7 Re:05;  
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWXF 6  
 DB 1 WXXWXF 6  
 RESULT 6  
 AAB01505  
 ID AAB01505 standard; peptide; 6 AA.  
 XX  
 AC AAB01505;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DE Peptide which binds to transcription factor E2F-1 DNA binding domain.  
 XX  
 KW DNA binding; transcription factor; E2F, E2F-1; cell cycle; DP-1;  
 KW activation; transcription; apoptosis; proliferative disorder;  
 KW psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 FH Key Location/Qualifiers  
 FT Misc-difference 2 /note= "Any amino acid"  
 FT Misc-difference 2 /note= "Any amino acid"  
 FT Misc-difference 2 /note= "Any amino acid"  
 FT Misc-difference 2 /note= "Any amino acid"

```

FT cancer
XX
XX
PS Example: Page 26; 42pp; English.
XX
CC Peptides which bind to the DNA binding domain of transcription
CC factor E2F and inhibit cell cycle progression may be useful as
CC research agents to investigate the interaction between E2F and DP-1,
CC or the activation of transcription by E2F 1/DP 1 heterodimers. They
CC may also be used for inducing apoptosis and/or cell cycle arrest in
CC a cell, particularly for treatment of cancer or other proliferative
CC disorders such as psoriasis and testecrosis.
XX
XX
SQ Sequence 6 AA;
XX
XX
Query Match 100.0%; Score 6; DB 21; Length 6;
Best Local Similarity 50.0%; Prod No 7 00000;
Matched 2; Conservation 2; Mismatch 0; Indels 0; Gaps 0;
XX
OY 1 WXXWVF 6
XX
DB 1 WAWWVF 6
XX
XX
RESULT 7
AAB01506
ID AAB01506 standard; Page 189; 6 AA
XX
AC AAE12181.
XX
XX
DT 08-NOV-2000 (first entry)
XX
XX
DE Peptide which binds to transcription factor E2F 1 DNA binding domain.
XX
XX
FW DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1,
XX
XX
KW activation, transcription, apoptosis, proliferative disorder;
XX
XX
XX
OS Synthetic.
XX
XX
PN WC020044771-A1.
XX
XX
PD 03-AUG-2000
XX
XX
PF 26-JAN-2000; 2006MO GP00027.
XX
XX
PR 26-JAN 1999; 993P 0001710
XX
XX
PA (PROL-) PROLIFIX LTD.
XX
XX
PI Mueller R, Kontermann RE, Montignani S;
XX
XX
PS WFI 2000-022807/4a
XX
XX
PT Peptides binding to the DNA binding domain of transcription factor
XX
XX
PT and inhibiting cell cycle progression, useful for the treatment of
XX
XX
PT cancer
XX
XX
PS Example: Page 26; 42pp; English.
XX
XX
CC Peptides which bind to the DNA binding domain of transcription
XX
XX
CC factor E2F and inhibit cell cycle progression may be useful as
XX
XX
CC research agents to investigate the interaction between E2F and DP 1,
XX
XX
CC or the activation of transcription by E2F 1/DP 1 heterodimers. They
XX
XX
CC may also be used for inducing apoptosis and/or cell cycle arrest in
XX
XX
CC a cell, particularly for treatment of cancer or other proliferative
XX
XX
CC disorders such as psoriasis and testecrosis.
XX
XX
SQ Sequence 6 AA;
XX
XX
Query Match 100.0%; Score 6; DB 21; Length 6;
Best Local Similarity 50.0%; Prod No 7 00000;
Matched 2; Conservation 2; Mismatch 0; Indels 0; Gaps 0;

```

```

OY 1 WXXWVF 6
XX
DB 1 WAWWVF 6
XX
XX
RESULT 8
AAB01508
ID AAB01508 standard; Page 189; 6 AA
XX
AC AAE12181.
XX
XX
DT 08-NOV-2000 (first entry)
XX
XX
DE Peptide which binds to transcription factor E2F 1 DNA binding domain.
XX
XX
FW DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1,
XX
XX
KW activation, transcription, apoptosis, proliferative disorder;
XX
XX
XX
OS Synthetic.
XX
XX
PN WC020044771-A1.
XX
XX
PD 03-AUG-2000.
XX
XX
PF 26-JAN 2000; 2006MO GP00027.
XX
XX
PR 26-JAN 1999; 993P 0001710.
XX
XX
PA (PROL-) PROLIFIX LTD.
XX
XX
PI Mueller R, Kontermann RE, Montignani S;
XX
XX
PS WFI 2000-022807/4a
XX
XX
PT Peptides binding to the DNA binding domain of transcription factor E2F
XX
XX
PT and inhibiting cell cycle progression, useful for the treatment of
XX
XX
PT cancer
XX
XX
PS Example: Page 26; 42pp; English.
XX
XX
CC Peptides which bind to the DNA binding domain of transcription
XX
XX
CC factor E2F and inhibit cell cycle progression may be useful as
XX
XX
CC research agents to investigate the interaction between E2F and DP-1,
XX
XX
CC or the activation of transcription by E2F 1/DP 1 heterodimers. They
XX
XX
CC may also be used for inducing apoptosis and/or cell cycle arrest in
XX
XX
CC a cell, particularly for treatment of cancer or other proliferative
XX
XX
CC disorders such as psoriasis and testecrosis.
XX
XX
SQ Sequence 6 AA;
XX
XX
Query Match 100.0%; Score 6; DB 21; Length 6;
Best Local Similarity 50.0%; Prod No 7 00000;
Matched 2; Conservation 2; Mismatch 0; Indels 0; Gaps 0;
XX
OY 1 WXXWVF 6
XX
DB 1 WAWWVF 6
XX
XX
RESULT 9
AAB01509
ID AAB01509 standard; Page 189; 7 AA
XX
AC AAE12181.
XX
XX
DT 18-DEC-2001 (first entry)
XX
XX
DE Polyglutamine repeat domain binding peptide, QP1-M7PD.
XX
XX
FW Polyglutamine repeat domain, Huntington, atrophin, ataxin
XX
XX
FW null, polyglutamine, Huntington, atrophin, ataxin,
XX
XX
XX
KW Huntington's disease, degenerative pallidum, atrophin, DRPLA;

```







stimulating factor (G-CSF a haematopoietic growth factor and cytokine that stimulates neutrophil proliferation and differentiation), including evaluation of many factors thought to influence, and be influenced by, production of white blood cells, in the development of compounds that bind to G-CSF, as reagents for detecting G-CSF receptor or related receptor on living cells, fixed cells, in biological fluid, in tissue homogenates or in purified natural biological materials, in situ staining, fluorescence-activated cell sorting (FACS), Western blotting or enzyme-linked immunosorbent assay (ELISA), in receptor purification or in purifying cells expressing G-CSF on the cell surface (or inside permeabilised cells) as a commercial research reagent for various medical and diagnostic uses or to treat a disease that would benefit from the ability to of a compound to mimic the effects of G-CSF in vivo.

The compounds bind specifically to G-CSF and allow for studies of biological activities mediated by the receptor and for the treatment of diseases, disorders and conditions that would benefit from activating or inactivating G-CSF. The present sequence is a G-CSF binding peptide of the invention.

Sequence 9 AA:

Query Match 100.0%; Score 6; DB 23; Length 9;  
Best Local Similarity 50.0%; Prod No 74479;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

OY 1 WXXXWF 6  
|:|:|  
DB 4 WWCWNP 9

RESULT 14  
AAW57400  
ID AAW57400 standard; Protein; 10 AA.  
XX  
AC AAW57400:  
XX  
DT 21-MAR-1995 (first entry)  
XX  
DE Peptide for treating diseases related to anti-DNA antibodies  
XX  
KW Carrier; absorbing agent; treatment; anti-DNA antibody; immune system  
XX  
OS Synthetic  
XX  
PN JPO6192290-A.  
XX  
PD 12-OTL-1994.  
XX  
PF 18-MAN-1994; 943P-0006098.  
XX  
PP 30-SEP-1992; 943P-001921.  
XX  
PA (KIPAS) KIPAPAY CO LTD.  
XX  
DR WPI; 1994-260519/32.  
XX  
PT A peptide and in absorbing agent; peptide; by immobilising it on a carrier - useful for treatment of diseases related to anti-DNA antibodies and immune complexes  
XX  
PS Disclosure; Page 12; 14pp; Japanese.  
XX  
CC The sequences given in AAW57396, 413 are peptides which are all covered by the claimed generic formula.  
CC H-X-(A-B)n-Y-Z  
CC A - Trp, Phe or a peptide fragment consisting of 2 residues;  
CC B = Trp, Phe, Asn or Glu;  
CC X and Y = a bond or Asp, Glu, Arg, Lys, His or a peptide fragment consisting of 2 to 10 residues, provided that at least one of X or Y are present;  
CC Z - OH or NH<sub>2</sub>; and  
CC n = 2-5.  
CC These peptides may be immobilised on a carrier in the preparation of an

absorbing agent which may be used in the treatment of diseases related to anti-DNA antibodies and/or immune complex.

Sequence 10 AA:

Query Match 100.0%; Score 6; DB 15; Length 10;  
Best Local Similarity 50.0%; Prod No 944792;  
Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

OY 1 WXXXWF 6  
|:|:|  
DB 3 WWCWNP 8

RESULT 15  
AAW62149  
ID AAW62149 standard; peptide; 11 AA.  
XX  
AC AAW62149;  
XX  
DT 16-SEP-1998 (first entry)  
XX  
DE Haemophilus influenzae tyrosine (TNA synthetase binding) peptide 22.  
XX  
KW Identification; ligand; biological activity; target-binding;  
XX drug screening; library; inhibitory ligand.  
XX  
OS Synthetic.  
XX  
OS Haemophilus influenzae.  
XX  
PN WO9819162-A1.  
XX  
PD 07-MAY-1998.  
XX  
PF 31-OCT-1997; 97WO-US19638.  
XX  
PP 31-OCT-1996; 96US-0740671.  
XX  
FA (HIVMA) NOVAMON PHARM CORP.  
XX  
TI Fowlkes CW, Fiedlinger JA, Hyde L et al; J Biol Chem; 1998; 273:89/24.  
XX  
DR WPI; 1998-272389/24.  
XX  
PT Identifying ligands which mediate biological activity of a protein  
XX by identifying target-binding ligands and screening a library for  
XX ligands which inhibit target-binding ligand mediated activity  
XX  
PS Example 5; Page 100; 143pp; English.  
XX  
CC A method has been developed for identifying a ligand which mediates the biological activity of a target protein (T) by inhibiting the binding of (T) to a binding partner. The method comprises: (a) screening a first combinatorial library comprising first member ligands for binding to the target-binding ligands (TBLs), to identifying one or more (PBLs); (b) screening a second library comprising second member ligands for the ability to inhibit the binding of (a) or (b) of the PBLs to the target protein, and so obtaining one or more inhibitory ligands; and (c) determining which of the inhibitory ligands can mediate a biological activity of the target protein. The present sequence represents a potential binding peptide for Haemophilus influenzae tyrosine (TNA synthetase from an example of the present invention. The method can be used for identifying drugs which can mediate the biological activity of a target protein. It can be used to identify the biological activity of a target protein whose biological function is not known and perhaps cannot be determined directly. The method can also be used to identify new inhibitory ligands of specific target proteins. The method provides high throughput screens which are essentially identical for similar and dissimilar targets, bypassing the need to develop distinct assays for biochemically diverse targets.

Sequence 11 AA:

Query Match 100.0% Score 67 14 19 Length 11;  
 Best Local Similarity 10.0% Freq 2 2 2 2 2  
 Matches 31 Conservative 31 Mismatches 0 Indels 0 Gaps 0  
 2Y 1 XXXXXF 6  
 1111  
 2b 5 WPDWGF 10

Search completed: February 20, 2003, 11:06:32  
 Job time : 30.313 secs









GenCore version 5.1.3  
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OM protein - protein search, using sw model

Run on: February 20, 2003, 10:41:55 / Search time 6.6667 seconds  
(without alignments)  
37 329 Million CPU updates/sec

Title: US-09-912-414-11

Perfect score: 38

Sequence: 1 WXXWHF 6

Scoring table: BLOSUM62DX

Gapop: 10.0, Gapext: 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 2000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database: SwissProt\_40.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being listed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	38	100.0	358	MTCL_CAUCR	O45971 Caulobacter
2	38	100.0	376	MTCL_SHIME	O30564 Rhizobium
3	38	100.0	377	MTCL_HYDRA	O30570 Brucella ab
4	38	100.0	424	FMCC_SOYBA	P48628 Glycine max
5	38	100.0	439	Y412_APAHA	O04658 Arabidopsis
6	38	100.0	443	FMCC_PPANA	P48627 Brassica na
7	38	100.0	448	FMCC_AARAH	P48312 Arabidopsis
8	38	100.0	479	CATA_PSEPI	O59714 Pseudomonas
9	38	100.0	504	S1F1_YEAST	O12460 Saccharomyc
10	38	100.0	511	NOPS_YEAST	O12460 Saccharomyc
11	38	100.0	570	NOPS_HUMAN	O21282 Homo sapien
12	38	100.0	574	NOPS_PAT	O21282 Homo sapien
13	38	100.0	540	Y412_EH13	P48628 Glycine max
14	38	100.0	718	PLISA_CARL	O21282 Homo sapien
15	35	92.1	140	LYSR_PROKE	O21282 Homo sapien
16	35	92.1	148	LYSR_PROKE	O21282 Homo sapien
17	35	92.1	140	LYSR_PROKE	O21282 Homo sapien
18	35	92.1	140	LYSR_PROKE	O21282 Homo sapien
19	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
20	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
21	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
22	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
23	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
24	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
25	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
26	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
27	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
28	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
29	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
30	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
31	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
32	35	92.1	142	LYSR_PROKE	O21282 Homo sapien
33	35	92.1	142	LYSR_PROKE	O21282 Homo sapien

34	32	84.2	85	1	VG22_BPM45	O02388 Mycobacteri
35	32	84.2	103	1	EGED_MYCIN	O05440 Mycobacteri
36	32	84.2	105	1	MGIM_HUMAN	O05179 Homo sapien
37	32	84.2	108	1	MGIM_POVYN	O05179 Homo sapien
38	32	84.2	111	1	VFX_HV2EP	O21282 Homo sapien
39	32	84.2	111	1	VFXM_SALTI	O21282 Homo sapien
40	32	84.2	111	1	VFXM_SALTY	O21282 Homo sapien
41	32	84.2	112	1	VFX_HV2CA	O21282 Homo sapien
42	32	84.2	112	1	VFX_HV2DI	O21282 Homo sapien
43	32	84.2	112	1	VFX_HV2SI	O21282 Homo sapien
44	32	84.2	112	1	VFX_HV2SI	O21282 Homo sapien
45	32	84.2	112	1	VFX_HV2SI	O21282 Homo sapien

## ALIGNMENTS

RESULT 1	ID	MTCL_CAUCR	STANDARD	PPT	358 AA
AC	O45971	MTCL_CAUCR	STANDARD	PPT	358 AA
DT	30-MAY-2000 (Rel. 39, Created)				
DT	16-OCT-2001 (Rel. 40, Last sequence update)				
DT	16-OCT-2001 (Rel. 40, Last annotation update)				
DE	Modification methylase CcmI (EC 2.1.1.72) (Adenine-specific				
DE	methyltransferase CcmI) (M.CcmI).				
GN	COPM OR CCM OR CCM378.				
OS	Caulobacter crescentus.				
OC	Bacteria; Proteobacteria; alpha subdivision; Caulobacter group;				
OC	Caulobacter.				
OX	NCBI TaxID=155892;				
PN	[1]				
FE	SEQUENCE FROM N.A.				
RC	STRAIN=CBEN / NA1000;				
PA	MEDLINE=94118203, PubMed=8289276;				
PA	Wiegler G, Marczynski G, Shapiro L;				
PT	"A Caulobacter DNA methyltransferase that functions only in the				
PT	predivisional cell";				
FT	J. Mol. Biol. 235:472-485(1994).				
RN	[2]				
RP	SEQUENCE FROM N.A.				
PC	STRAIN=ATCC 19089 / CB15;				
PX	MEDLINE=9173648, PubMed=11250647;				
PA	Nierman W.C., Foldvary T.V., Laub M.T., Paulsen I.T., Nelson K.E.,				
PA	Eisen T., Heidelberg J.F., Alley W.P., Ohta N., Maitav Y.P.,				
PA	Fotocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely H.,				
PA	Neely R.T., Deason P.J., Durkin A.S., Gilm M.B., Hart D.H.,				
PA	Pollock D.P., Tran K., Graven M.B., Khorat H., Shetty T., Berry K.,				
PA	Heidelberg J.F., Wolf A., Vamathevan J., Finkelstein M., White G.,				
PA	Salzberg S.L., Venter A.C., Shapiro L., Fraser C.M.);				
PT	"Complete genome sequence of Caulobacter crescentus";				
PT	Proc. Natl. Acad. Sci. U.S.A. 98:417-421(2001)				
CC	1. FUNCTION: THIS METHYLASE RECOGNIZES THE TGAATTA SEQUENCE				
CC	SAME AND CUES SPECIFIC METHYLATION AT A TGAATTA SEQUENCE.				
CC	FROM METHYLATED METHYLATION HAS IMPROVED CELLULAR FUNCTIONS.				
CC	APPEARS TO CONTRIBUTE TO THE ACTIVATE CELL-CELL INTERACTION OF DNA				
CC	REPLICATION AND CELLULAR MORPHOLOGY.				
CC	2. CATALYTIC ACTIVITY: S-adenosyl-L-methionine + DNA (adenine - S-				
CC	adenosyl-L-homocysteine + DNA 6-methylaminopurine.				
CC	3. This Swiss-Prot entry is deprecated. It is replaced through a collaboration				
CC	between the Swiss Institute of Bioinformatics and the EMBL database on its				
CC	the European Bioinformatics Institute. There are no restrictions on its				
CC	use by non-profit institutions as long as its content is in no way				
CC	modified and this statement is not removed. Usage by and for commercial				
CC	enterprises requires a license agreement (see <a href="http://www.ebi.ac.uk/submit/">http://www.ebi.ac.uk/submit/</a>				
CC	or send an email to <a href="mailto:license@ebi-sib.ch">license@ebi-sib.ch</a> ).				
CC	EMBL: U01032; AAA18913.1;				
DR	EMBL: A000571; NA022365.1;				
DR	HSEF: P11473; 1800;				
DR	PERASE; 2539; M.CcmI.				
DR	TIGR: CC0378; -				









CC THE SECOND ENTRY IN THE PROPERTIES OF 1.3 AND 1.4 FATTY  
CC ACTS, IMPROVED PROPERTIES OF PLANT MEMBRANES. IT IS THE  
CC THE FATTY ACID AS AN ELECTRON DONOR AND ACT AS FATTY ACIDS  
CC ESTERIFIED TO GALACTOLIPIDS, SULFOLIPIDS AND PHOSPHATIDYLCHOLINE  
CC PATHWAY POLYUNSATURATED FATTY ACID BIOSYNTHESIS  
CC SUPERFICIAL LOCATION: CHLOROPLAST MEMBRANE (P-PROTEIN)  
CC -1- DEVELOPMENTAL STAGE: HIGHEST LEVELS FOUND IN EXPANDING LEAVES  
CC -1- DOMAIN: THE HIGHEST LOW DOMAINS MAY CONTAIN THE ACTIVE SITE  
CC AND/OR BE INVOLVED IN METAL ION BINDING.  
CC -1- SIMILARITY: BELONGS TO THE FATTY ACID DESATURASE FAMILY.  
CC  
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CC or send an email to [license@sib-sib.ch](mailto:license@sib-sib.ch))  
CC  
CC EMBL; U09503; AAA92800.1; -  
CC EMBL; AL022198; CA18198.1; -  
CC EMBL; AL161578; CAB79813.1; -  
CC EMBL; AY045623; AAK72979.1; -  
CC EMBL; AY058078; AAL24186.1; -  
CC EMBL; AY058078; AAL24240.1; -  
CC InterPro; IPR001225; FA desaturase.  
CC Pfam; PF00487; FA desaturase; 1.  
CC ProDom; PD001081; FA desaturase; 1.  
CC OxiReductase; Fatty acid biosynthesis, Chloroplast, Membrane,  
CC TRANSIT peptide.  
CC TRANSIT 1 69 CHLOROPLAST (BY SIMILARITY).  
CC CHAIN 70 448 OMEGA-6 FATTY ACID DESATURASE.  
CC FT DOMAIN 171 175 HISTIDINE BOX-1.  
CC FT DOMAIN 207 211 HISTIDINE BOX-2.  
CC FT DOMAIN 367 371 HISTIDINE BOX-3.  
CC SEQUENCE 448 AA, 5135 MW, C3AC72FE23FE2281 CRC64,  
CC  
CC Query Match 100.0%; Score 38; DB 1; Length 448;  
CC Best Local Similarity 66.7%; Freq. No. 61;  
CC Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
CC  
CC 1 WXXWHF 6  
CC  
CC 257 WXXWHF 262  
CC  
CC RESULT 8  
CC CATA\_PSEPU STANDARD; PRT; 479 AA.  
CC ID CATA\_PSEPU STANDARD; PRT; 479 AA.  
CC AC Q59714;  
CC DT 30-MAY-2000 (rel. 39, Created)  
CC DT 30-MAY-2000 (rel. 39, Last sequence update)  
CC DT 30-MAY-2000 (rel. 39, Last annotation update)  
CC DE Catalase (EC 1.11.1.6).  
CC GN KATA-OP CATA.  
CC OS Pseudomonas putida  
CC OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;  
CC Pseudomonas  
CC NCBI\_TaxID=303;  
CC RN [1]\_TaxID=303;  
CC RP SEQUENCE FROM N.A.  
CC RC STRAIN=Corvallis;  
CC RX MEDLINE=9801991; PubMed=9358059;  
CC RA Kim Y C, Miller C D, Anderson A J;  
CC RT Identification of adjacent genes encoding the major catalase and a  
CC RT bacterioferritin from the plant-beneficial bacterium Pseudomonas  
CC RT putida";  
CC RL Gene 1991219-224(1997)  
CC  
CC FUNCTION REGULATOR OF HYDROGEN PEROXIDE IN WATER AND HYDROGEN PEROXIDE  
CC TO PROTECT CELLS FROM THE TOXIC EFFECTS OF HYDROGEN PEROXIDE  
CC -1- CATALYTIC ACTIVITY: 2 H2O2(O2) -> O2 + 2 H2O.  
CC -1- COFACTOR: HEME GROUP.  
CC -1- ENZYME REGULATION: ACTIVATED BY PEROXIDE.

CC -1- SIMILARITY: BELONGS TO THE CATALASE FAMILY.  
CC  
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CC  
CC EMBL; U63511; AAB88219.1; -  
CC HSSP; F4231, COAE.  
CC InterPro; IPR002226; Catalase.  
CC Pfam; PF00199; Catalase; 1.  
CC EPRINTS; PR00067; CATALASE.  
CC PRODOM; PR000510; Catalase; 1.  
CC DP PROSITE; PS00437; CATALASE; 1; 1.  
CC DP PROSITE; PS00438; CATALASE; 2; 1.  
CC KW Oxidoreductase, Peroxidase, Iron, Heme, Hydrogen peroxide,  
CC ACT SITE 53 53 BY SIMILARITY.  
CC FT ACT SITE 126 126 PROXIMAL HEME LIGAND (BY SIMILARITY).  
CC FT BINDING 336 336  
CC SEQUENCE 479 AA, 53381 MW, EEE3C8D67778571 CRC64,  
CC  
CC Query Match 100.0%; Score 38; DB 1; Length 479;  
CC Best Local Similarity 66.7%; Freq. No. 64;  
CC Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
CC  
CC 1 WXXWHF 6  
CC  
CC 209 WXXWHF 214  
CC  
CC RESULT 9  
CC SIK1\_YEAST STANDARD; PRT; 504 AA.  
CC ID SIK1\_YEAST STANDARD; PRT; 504 AA.  
CC AC Q12460;  
CC DT 01-NOV-1997 (rel. 35, Created)  
CC DT 01-NOV-1997 (rel. 35, Last sequence update)  
CC DT 15-JUN-2000 (rel. 41, Last annotation update)  
CC DE SIK1 protein (Nucleolar protein NOP56).  
CC GN SIK1 OR NOP56 OR YLP197W OR L0167.9.  
CC OS Saccharomyces cerevisiae (Baker's yeast).  
CC CC Eukaryota, Fungi; Ascomycota, Saccharomycetes;  
CC CC Saccharomycetales; Saccharomycetaceae; Saccharomyces.  
CC NCBI\_TaxID=4932;  
CC RN [1]\_TaxID=4932;  
CC RP SEQUENCE FROM N.A.  
CC RC STRAIN=S288c / YPH1;  
CC RX MEDLINE=96040178; PubMed=7547500;  
CC MOIRIN F J, Downs J A, Shadgrass A M, Gilmore T D;  
CC "Genetic analysis of growth inhibition by GNA1-1 kappa P-199 in  
CC RT Saccharomyces cerevisiae";  
CC RL Cell Growth Differ. 6:789-798(1995).  
CC RN [2]  
CC RP SEQUENCE FROM N.A.  
CC RC STRAIN=S288c / AB972;  
CC RA Johnson M, Andrews S, Printman F, Cooper J, Ding H, Du Z,  
CC RA Favello A, Patton L, Gattling S, Grewer T, Kistner J, Kozuka T,  
CC RA Hallsworth K, Hawkins J, Hillier L, Jermolov, Johnson D,  
CC RA Johnston L, Langston V, Lattelle F, Mardis E, Meneses S,  
CC RA Miller N, Nham M, Pauley A, Peluso D, Ritten L, Riles D,  
CC RA Tach S, Trevasis F, Vaudin A, Vignati D, Wilcox L, Wilson R,  
CC RA Woldman P, Waterston R;  
CC RL Submitted (SEP-1994) to the EMBL/GenBank/DBD databases.  
CC RN [3]  
CC RP CHARACTERIZATION AND MUTAGENESIS.  
CC RX MEDLINE=98038777; PubMed=9772949;  
CC RA Jachler T, Bajcs T, Tolstevy D, Hurt E;  
CC RT "Nucleolar P2E/D repeat proteins NOP56 and NOP57 interact with NOP1p  
CC RT and are required for ribosome biogenesis";  
CC RL Mol Cell Biol 17:7084-7094(1997)  
CC  
CC FUNCTION REGULATOR FOR THE BIOSYNTHESIS OF NUCLEOTIDES.



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DR EMBL: AF123534; AAD27610.1;  
 DR EMBL: AF261608; AAF3134.1;  
 DR EMBL: AL117554; CAB5598.1;  
 DR EMBL: AF161469; AAF3094.1;  
 DR EMBL: AF161469; AAF3094.1;  
 DR SWISS-EXPAGE; GYEX3; HUMAN.  
 DR InterPro; IPR026687; NIP.  
 DR Pfam; PF01798; NOP; 1.  
 DR ProDom; PD004104; NOP; 1.  
 KW Ribosome biogenesis; Nuclear protein.

FT CONFLICT 2 2 L->M (IN REF. 3).  
 FT CONFLICT 129 129 G->V (IN REF. 3).  
 FT CONFLICT 129 129 LTVQVQGVQVDFEYVAGAV -> VHTASVETZALPRLC.  
 FT CONFLICT 129 129 CO (IN REF. 4).  
 FT CONFLICT 235 260 RAAAEISMSTVESEEDICNIIHICQ -> EGGRPIHICSP.  
 FT CONFLICT 260 260 GPPPIYQVSASLHP (IN REF. 4).  
 FT CONFLICT 444 447 M->V (IN REF. 3).  
 FT CONFLICT 444 447 PRLEQ -> TQNT (IN REF. 3).  
 SQ SEQUENCE 579 AA; 6479 MW; 370930FEF9A56C PROQA;

Query Match 100.0%; Score 39; DP 1; Length 579;  
 Best Local Similarity 66.7%; Pred. No. 70;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 1 WXXWHF 6  
 DB 186 WYGMHF 191

RESULT 12  
 NOPS\_RAT STANDARD; PRT; 534 AA.  
 ID NOPS\_RAT  
 AC Q9QZ65; O88525;  
 DT 16-OCT-2001 (Rel. 40; Created)  
 DT 16-OCT-2001 (Rel. 40; Last sequence update)  
 DT 15-JUN-2000 (Rel. 41; Last annotation update)  
 DE Nucleolar protein NOPS (Nucleolar protein 5) (Nopp140 associated  
 DE protein).  
 GN NOPS\_OP NAB65.  
 OS Rattus norvegicus (Rat).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eumetazoa;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus  
 OC NCB1\_TaxID 10116.  
 RN (1)  
 RP SEQUENCE FROM N.A. SEQUENCE OF 1.23. AND INTERACTION WITH NOPS1  
 RA MEDLINE 2014370; PubMed 10472015;  
 RA Yama Y, Isaac C, Wang C, Dragan P, Paganis V, Moller H T;  
 RT "Conserved composition of mammalian box H/A/A and box C/C/C motif  
 RT nucleolar ribonucleoprotein particles and their interaction with the  
 RT common factor Nopp140.";  
 RT Mol. Biol. Cell 11:567-577(2000).  
 RI (2)  
 PP SEQUENCE OF 1-461 FROM N.A.  
 RA Hutton D, Gray J C;  
 RI Submitted (JUN 1998) to the EMBL/GenBank/DBP databases.  
 CC 1- FUNCTION: REGULATOR FOR AND PROSOMAL SEGMENT BIOGENESIS (By  
 CC similarity).  
 CC 1- SUBUNIT: Interacts with Ndc1/Nopp140.  
 CC 1- SUBCELLULAR LOCATION: Nuclear; nucleolar.  
 CC 1- SIMILARITY: BELONGS TO THE NOPS/NOP56 FAMILY.

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DR EMBL: AF193377; AAF05769.1;  
 DR EMBL: AF069782; AAC23535.1;  
 DR InterPro; IPR026687; NOP.  
 DR Pfam; PF01798; NOP; 1.  
 DR ProDom; PD004104; NOP; 1.  
 KW Ribosome biogenesis; Nuclear protein.  
 FT CONFLICT 396 396 E->K (IN REF. 2).  
 FT CONFLICT 459 459 A->K (IN REF. 2).  
 SQ SEQUENCE 534 AA; 60070 MW; 4B3585FA1E67779 CRC64;

Query Match 100.0%; Score 39; DP 1; Length 534;  
 Best Local Similarity 66.7%; Pred. No. 71;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 1 WXXWHF 6  
 DB 186 WYGMHF 191

RESULT 13  
 Y4CD\_RHISN STANDARD; PRT; 619 AA.  
 ID Y4CD\_RHISN  
 AC P55386;  
 DT 01-NOV-1997 (Rel. 35; Created)  
 DT 01-NOV-1997 (Rel. 35; Last sequence update)  
 DT 01-NOV-1997 (Rel. 35; Last annotation update)  
 DE Hypothetical 71.6 kDa protein Y4CD.  
 GN Y4CD.  
 OS Rhizobium sp. (strain NCR234).  
 OS Plasmid sym PNGP234a.  
 OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae Group;  
 OC Rhizobiaceae; Rhizobium  
 OC NCB1\_TaxID=394;  
 RN (1)  
 RP SEQUENCE FROM N.A.  
 RA MEDLINE 97205956; PubMed 9161424;  
 RA Freiberg C A, Pailley P, Pailley A, Proughon W J, Rosenthal A,  
 RA Porret X,  
 RT "Molecular basis of symbiosis between Rhizobium and legumes.";  
 RT Nature 387 394-401(1997).  
 RI (2)  
 CC 1 SIMILARITY: NONE OBVIOUS.

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 CC or send an email to [license@sib-sib.ch](mailto:license@sib-sib.ch))

DR EMBL: AF090667; AAB91634.1;  
 DR InterPro; IPR026157; TR\_Jdomain.  
 DR SMART; SMART5; TIR; 1.  
 KW Hypothetical protein; Plasmid.  
 SQ SEQUENCE 646 AA; 71604 MW; 81FE0A0A0A0E CF164;

Query Match 100.0%; Score 38; DP 1; Length 640;  
 Best Local Similarity 66.7%; Pred. No. 84;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 1 WXXWHF 6  
 DB 521 WPFWHF 526

RESULT 14  
 PLSB\_CAEEL STANDARD; PRT; 718 AA.  
 ID PLSB\_CAEEL  
 AC Q22949;  
 DT 15-DEC-1998 (Rel. 37; Created)  
 DT 15-DEC-1998 (Rel. 37; Last sequence update)



DR PIR; S12726; S12726;  
 DR HSP; P00695; LZ5;  
 DR Flyase; FBgn001120; LysA.  
 DR Flyase; FBgn004426; LysC.  
 DR Flyase; FBgn004427; LysD.  
 DR Interpio; IPRO01916; GH\_22.  
 DR Pfam; PF00062; Lys; 1.  
 DR PRINTS; PR00135; LYZLACT.  
 DR SMART; SM00263; LYZ1; 1.  
 DR pPocite; P00118; LACTALDEHYD\_LYSOZYME; 1.  
 KW Hydrolase; Glycosidase; Bacteriolytic enzyme; Signal;  
 KW Multigene family.  
 FT SIGNAL 1 18 BY SIMILARITY.  
 FT CHAIN 19 140 LYSOZYME A/C/D.  
 FT DISULFID 24 139 BY SIMILARITY.  
 FT DISULFID 45 129 BY SIMILARITY.  
 FT DISULFID 80 96 BY SIMILARITY.  
 FT DISULFID 92 110 BY SIMILARITY.  
 FT ACT\_SITE 50 50 BY SIMILARITY.  
 FT ACT\_SITE 68 68 BY SIMILARITY.  
 FT CONFLICT 41 42 AP -> NK (IN REF. 1)  
 SO SEQUENCE 140 AA; 15615 MW; 75024C6F5DF90; CRC64;

Query Match 92.18; Score 35; DB 1; Length 140;  
 Best Local Similarity 50.0%; Pred. No. 56;  
 Matches 3; Conservative 1; Mismatches 0; Indels 0;  
 Gaps 0;

OY 1 WXXWHF 6  
 DB 123 WSTWHY 128

Search completed: February 20, 2003, 10:44:03  
 Job time : 7.66667 secs

1. The first part of the document discusses the importance of maintaining accurate records of all transactions and activities. It emphasizes that proper record-keeping is essential for transparency and accountability, particularly in financial matters. The text outlines various methods for organizing and storing data, including digital databases and physical filing systems.

2. The second section focuses on the role of communication in project management. It highlights the need for clear, concise, and timely communication between team members and stakeholders. The author provides several strategies for effective communication, such as regular meetings, status reports, and the use of collaborative tools.

3. The third part of the document addresses the challenges of resource allocation. It discusses how to identify and prioritize tasks, allocate resources efficiently, and manage potential bottlenecks. The text includes practical advice on how to handle competing demands and ensure that all team members have the necessary resources to complete their work.

4. The final section discusses the importance of flexibility and adaptability in project management. It notes that projects often encounter unexpected changes and challenges, and that successful managers must be able to respond quickly and effectively to these changes. The author provides examples of how to adjust plans and strategies as needed to keep the project on track.



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## OM Protein Protein search using SW model

Run on February 20, 2003, 10:41:10, Search time: 15.413 seconds  
(without alignment)

52944 Million cell features/sec

Title: US-09-912-414-11

Perfect score: 38

Sequence: 1 WXXWHF 6

Database: RPTREMBL

Gapop: 10.0, Gapext: 0.5

Scanned: 671580 seqs, 26604715 residues

Total number of hits satisfying chosen parameters: 47544

Minimum DB seq length: 0

Maximum DB seq length: 260000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database:

1: SP\_TREMBL\_21:  
2: sp\_alchea:  
3: sp\_bacteria:  
4: sp\_fungi:  
5: sp\_human:  
6: sp\_invertebrate:  
7: sp\_mhc:  
8: sp\_mammal:  
9: sp\_mhc:  
10: sp\_mhc:  
11: sp\_mhc:  
12: sp\_mhc:  
13: sp\_mhc:  
14: sp\_mhc:  
15: sp\_mhc:  
16: sp\_mhc:  
17: sp\_mhc:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	38	100.0	147	Q91204	Q91204 Pasteurella
2	38	100.0	140	Q91205	Q91205 Pasteurella
3	38	100.0	201	Q91206	Q91206 Pasteurella
4	38	100.0	202	Q91207	Q91207 Pasteurella
5	38	100.0	213	Q91208	Q91208 Pasteurella
6	38	100.0	214	Q91209	Q91209 Pasteurella
7	38	100.0	215	Q91210	Q91210 Pasteurella
8	38	100.0	216	Q91211	Q91211 Pasteurella
9	38	100.0	217	Q91212	Q91212 Pasteurella
10	38	100.0	218	Q91213	Q91213 Pasteurella
11	38	100.0	219	Q91214	Q91214 Pasteurella
12	38	100.0	220	Q91215	Q91215 Pasteurella
13	38	100.0	221	Q91216	Q91216 Pasteurella
14	38	100.0	222	Q91217	Q91217 Pasteurella
15	38	100.0	223	Q91218	Q91218 Pasteurella
16	38	100.0	224	Q91219	Q91219 Pasteurella

17	38	100.0	381	Q91204	Q91204 Pasteurella
18	38	100.0	386	Q91205	Q91205 Pasteurella
19	38	100.0	394	Q91206	Q91206 Pasteurella
20	38	100.0	403	Q91207	Q91207 Pasteurella
21	38	100.0	415	Q91208	Q91208 Pasteurella
22	38	100.0	421	Q91209	Q91209 Pasteurella
23	38	100.0	423	Q91210	Q91210 Pasteurella
24	38	100.0	424	Q91211	Q91211 Pasteurella
25	38	100.0	424	Q91212	Q91212 Pasteurella
26	38	100.0	425	Q91213	Q91213 Pasteurella
27	38	100.0	425	Q91214	Q91214 Pasteurella
28	38	100.0	425	Q91215	Q91215 Pasteurella
29	38	100.0	425	Q91216	Q91216 Pasteurella
30	38	100.0	425	Q91217	Q91217 Pasteurella
31	38	100.0	425	Q91218	Q91218 Pasteurella
32	38	100.0	425	Q91219	Q91219 Pasteurella
33	38	100.0	425	Q91220	Q91220 Pasteurella
34	38	100.0	425	Q91221	Q91221 Pasteurella
35	38	100.0	425	Q91222	Q91222 Pasteurella
36	38	100.0	425	Q91223	Q91223 Pasteurella
37	38	100.0	425	Q91224	Q91224 Pasteurella
38	38	100.0	425	Q91225	Q91225 Pasteurella
39	38	100.0	425	Q91226	Q91226 Pasteurella
40	38	100.0	425	Q91227	Q91227 Pasteurella
41	38	100.0	425	Q91228	Q91228 Pasteurella
42	38	100.0	425	Q91229	Q91229 Pasteurella
43	38	100.0	425	Q91230	Q91230 Pasteurella
44	38	100.0	425	Q91231	Q91231 Pasteurella
45	38	100.0	425	Q91232	Q91232 Pasteurella

## ALIGNMENTS

RESULT 1	Q91204	PRELIMINARY:	PRT:	187 AA.
ID	Q91204			
AC	Q91204			
DT	01-JUN-2001 (TREMREL: 17, last sequence update)			
DT	01-JUN-2001 (TREMREL: 17, last sequence update)			
DT	01-MAR-2002 (TREMREL: 19, last annotation update)			
DE	Hypothetical protein PM135.			
GN	Pasteurella multocida.			
OS	Pasteurella multocida.			
OC	Bacteria; Proteobacteria; gamma subdivision; Pasteurellales;			
OC	Pasteurella.			
OX	NCBI_TaxID=747;			
RN	(1)			
RF	SEQUENCE FROM N.A.			
FC	STRAIN=EM70;			
PX	MEDLINE:2145866, PubMed:1124100;			
RA	May B.J., Zhang Q., Li L.L., Paustian M.L., Whitlam T.S., Kapur V.;			
RT	"Complete genomic sequence of Pasteurella multocida PM135";			
PL	Proc Natl Acad Sci U S A 98:1460-1465(2001);			
CC	1-1 SIMILARITY. RELINGS TO THE TET/ACR FAMILY OF TRANSCRIPTIONAL			
CC	REGULATORS.			
DP	EMBL: AE006154; AAF03219.1; -			
DR	InterPro: IPR01647; HTH_Tetr.			
DP	Pfam: PF0440; Tetr_1.			
DP	PRINTS: PR00455; HTHTR.			
KW	CNA-binding, Hypothetical protein; Transcription regulation;			
KW	Complete proteome.			
SQ	SEQUENCE 187 AA, 2025 MW, -38290ECAL111F D384;			

Query Match: 100.0%, Score: 32, E: 1.6, Length: 187;

Best Local Similarity: 66.7%, Pred. No.: 1.0e+00;

Matches: 4; Conservative: 0; Mismatches: 0; Indels: 0; Gaps: 0;

QY 1 WXXWHF 6

DB 89 WXXWHF 94

[illegible]

Query Match	100.00; Score 38; DB 10; Length 202;
Best Local Similarity 44.7%; Prod. No. 2e-02;	
Matching 4; Conservative 2; Miscellaneous 0; Indexes 0; Gaps 1;	

[illegible]

John W. I	1	3	John J.	28	16	John J.	23
Eastwood	Similarity	69.7%	Prod. No.	2.1e+02			
Marshall	1	Conservative	2	Misatches	0	Totals	0
						Gaps	0

Db 154 WVMWHF 169

RESULT 6  
09AXY2 PRELIMINARY, FFT, 216 AA.

AC 09AXY2;  
DT 01-JUN-2001 (TEMBLrel, 17, last sequence update)  
DT 01-JUN-2002 (TEMBLrel, 21, last annotation update)  
DE Omega 6 reductase (fragment).  
GN FAD6-BN-1.  
OS Brassica napus (rape).  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons; Rosidae;  
OC eucosids II; Brassicales; Brassicaceae; Brassica.  
OX NCBI\_TaxID=3708;

RA SEQUENCE FROM N.A.  
RC STRAIN=CV, STELLAR.  
RA Fourman M., Froger N., Brunel D.; Tools designing for a genetic map  
RT "Amplified consensus gene markers for mapping for a genetic map  
RL of Arabidopsis known-function genes in Brassica";  
RL Submitted (JAN-2000) to the EMBL/GenBank/DBS databases.  
DR EMBL; AF229391; AAK0663.1;  
DR InterPro: IPR001225; FA desaturase  
DR Pfam: PF00487; FA desaturase; 1.  
DR ProDom: PD001081; FA desaturase; 2.  
FT NON\_TER 1 1  
FT NON\_TER 216 216  
SQ SEQUENCE 216 AA; 2510 MW; EFEEEBEHBIEHIEA CPG64.

Query Match 100.0%; Score 19; DB 19; Length 216;  
Best Local Similarity 66.7%; Prod No. 2 1e+02;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
Db 154 WVMWHF 162

RESULT 7  
094K47 PRELIMINARY, PRT, 219 AA.

AC 094K47;  
DT 01-DEC-2001 (TEMBLrel, 19, last sequence update)  
DT 01-JUN-2002 (TEMBLrel, 21, last annotation update)  
DE Putative 6-phosphogluconolactonase  
OC Arabidopsis thaliana (mouse-ear cress).  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons;  
OC eucosids II; Brassicales; Brassicaceae; Arabidopsis.  
OX NCBI\_TaxID=3702;

RA SEQUENCE FROM N.A.  
RA Yamada K., Liu S.Y., Sakano H., Phan P.Y., Path T., Chung W.Y.,  
RA Goldsmith A.D., Lee J.M., Quach H.L., Tang C.C., Toriumi M., Yu G.,  
RA Bowers J., Carimato P., Chen H., Chey P., Hayashiroki Y., Ishida C.,  
RA Jones T., Kamiya A., Karlin-Newman G., Kawai T., Kim C., Kosuma R.,  
RA Lam P., Lin J., Meyers W.C., Miranda M., Narusaka M., Nguyen M.,  
RA Pal M.C., Sakurai T., Satoh M., Seki M., Shino P., Southwick A.,  
RA Shinkai Y., Davis P.W., Becker J.P., Theologis A.,  
RT "Full length cDNA of gene FGAH7.11/ATG54400 (GI:10179791)";  
RL Submitted (APR-2001) to the EMBL/GenBank/DBS databases  
CC "SIMILARITY SEARCHING 1 WD FILTER (SEE ABL DOMAIN)".  
DB EMBL; AF270305; AAK44120.1;  
DB TIGRPFAMS; TIGR01198; p01.1;  
KW Repeat; WD repeat.  
SQ SEQUENCE 219 AA; 24540 MW; 1904A77323434RA CPG64.

Query Match 100.0%; Score 39; DB 10; Length 219;

Best Local Similarity 66.7%; Prod No. 2 1e+02;  
Matches 4; Conservative 2; Mismatches 0; Indels 0;  
Db 72 WVMWHF 77

Query 1 WXXWHF 6

RESULT 8  
09AXY4 PRELIMINARY, FFT, 222 AA.

AC 09AXY4;  
DT 01-JUN-2001 (TEMBLrel, 17, last sequence update)  
DT 01-JUN-2001 (TEMBLrel, 17, last sequence update)  
DT 01-JUN-2002 (TEMBLrel, 21, last annotation update)  
DE Omega 6 reductase (fragment).  
GN FAD6-BO-1.  
OS Brassica oleracea (cauliflower).  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons;  
OC eucosids II; Brassicales; Brassicaceae; Brassica.  
OX NCBI\_TaxID=3712;

RA SEQUENCE FROM N.A.  
RC STRAIN=CV, RAPID CYCLING.  
RA Fourman M., Froger N., Brunel D.; Tools designing for a genetic map  
RT "Amplified consensus gene markers for mapping for a genetic map  
RL of Arabidopsis known-function genes in Brassica";  
RL Submitted (JAN-2000) to the EMBL/GenBank/DBS databases.  
DR EMBL; AF229399; AAK0661.1;  
DR InterPro: IPR001225; FA desaturase.  
DR Pfam: PF00487; FA desaturase; 1.  
DR ProDom: PD001081; FA desaturase; 2.  
FT NON\_TER 1 1  
FT NON\_TER 222 222  
SQ SEQUENCE 222 AA; 2601 MW; AAF1F4W4WPTAE CPG64;

Query Match 100.0%; Score 38; DB 10; Length 222;  
Best Local Similarity 66.7%; Prod No. 2 2e+02;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
Db 163 WVMWHF 168

Query 1 WXXWHF 6

RESULT 9  
053604 PRELIMINARY, FFT, 230 AA.

AC 053604;  
DT 01-JUN-1998 (TEMBLrel, 06, last sequence update)  
DT 01-JUN-1998 (TEMBLrel, 06, last sequence update)  
DT 01-MAR-2002 (TEMBLrel, 20, last annotation update)  
DE Hypothetical protein RV0059.  
GN FvG02 OF MTW10 22 OF MTW065.  
OS Mycobacterium tuberculosis.  
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;  
OC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.  
OX NCBI\_TaxID=1773;

RA SEQUENCE FROM N.A.  
RC STRAIN=H37RV,  
PX MEDLINE; 98325987; PubMed:9634230,  
RA Cole S.T., Broch R., Finkhull J., Guller T., Church C., Harris D.,  
RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,  
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,  
RA Davies R., Devlin K., Felwell T., Gentles E., Hamlin R., Holtgrave S.,  
RA Horsby T., Jagers K., Krogh A., McLean J., Mole S., Murphy D.,  
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,  
RA Rutter S., Seeger K., Skelton S., Squares S., Squares P.,  
RA Sulston J.E., Taylor K., Whithead S., Wallis J.G.;  
RT "Deciphering the biology of Mycobacterium tuberculosis from the  
complete genome sequence.";



Db 220 WBPWHF 225

## RESULT 13

ID Q8TRL3 PRELIMINARY FFT 293 AA

AC Q8TRL3

DT 01-JUN-2002 (TrEMBLrel 21, Created)

DT 01-JUN-2002 (TrEMBLrel 21, Last sequence update)

DT 01-JUN-2002 (TrEMBLrel 21, Last annotation update)

DE Hypothetical protein M11162.

GN M11162

OS Methanococcus acetivorans

OC Archaea; Euryarchaeota; Methanococci; Methanococciales;

OC Methanococcinaceae; Methanococcinae

OX NCBI\_TaxID=2214

RN (1)

RP SEQUENCE FROM N.A.

RC STRAIN 22A / ATCC 35967 / DSM 2034

RX MEDLINE=21929760; PubMed=1133238

RA Galian J.E., Musbaum C., Roy A., Fredrick M.G., Madenjian P.,

RA Fitzhugh M., Galvo S., Engels R., Smirnov S., Atwood D., Brown A.,

RA Allen N., Myler J., Stange-Thomann R., DeArlano P., Johnson P.,

RA Linco J., McKen P., Wierman P., Thomas J., Strick A., Ye W.,

RA Zimmer A., Barber P.D., Cann I., Graham D.E., Graham A.M.,

RA Hedderich P., Ingram-Smith C., Krieger H.C., Krzycki J.A.,

RA Leigh J.A., Li W., Liu J., Mukhopadhyay R., Reeve J.N., Smith K.,

RA Springer T.A., Tanyali A., White O., White P.H., de Marco R.C.,

RA Pritchett M., Sowers P.P., Swanson P.V., Zinder S.H., Zinder F.,

RA Metcalf W.W., Birren B.

PT "The genome of Methanococcus acetivorans reveals extensive metabolic

RT and physiological diversity."

RL Genome Res. 13:432-442(2003)

DR EMBL: AEO10783; AM04583.1

KM Hypothetical protein; Complete proteome

SQ SQUINTE 293 AA; 33721 MW; 44971/45949/46964 CPG64

Query March 100.0% Score 39, FR 17, Length 293

Best Local Similarity 66.7% Pred No 2e+02

Matches 4; Conservative 2; Mismatches 3; Indels 0; Gaps 0

QY 1 WXXWHF 6

Db 168 WSLWHF 173

RESULT 14

ID Q91065 PRELIMINARY FFT 293 AA

AC Q91065

DT 01-MAR-2001 (TrEMBLrel 16, Created)

DT 01-MAR-2001 (TrEMBLrel 16, Last sequence update)

DT 01-JUN-2002 (TrEMBLrel 21, Last annotation update)

DE Hypothetical protein PA2778.

GN PA2778

OS Pseudomonas aeruginosa

OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;

OC Pseudomonas

OX NCBI\_TaxID=287

RN (1)

RP SEQUENCE FROM N.A.

RC STRAIN-ATCC 15692 / PA01

RX MEDLINE=20437337; PubMed=10984043

RA Sawyer C.K., Pham X.-O.T., Erwin A.L., Mizoguchi S.D., Martner P.,

RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou Y.,

RA Garber P.L., Coltray L., Tolentino E., Westbrock-Wadman S., Yuan Y.,

RA Brody L.L., Coltray S.N., Folger P.P., Vas A., Barry J., Lim P.M.,

RA Smith P.A., Spencer D.H., Wong G.K.W., Wu Z., Paulsen I.T.,

RA Reizer J., Sailer M.H., Hancock P.B.W., Lory S., Olson M.V.,

RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an

RT opportunistic pathogen."

RL Nature 406:59-64(2000)

DR EMBL: AEO44705; AA006166.1

DR InterPro: IPR005074; Peptidase\_C39.

DR InterPro: IPR001440; TPR

DR Pfam: PF03412; Peptidase\_C39, 1.

KW Hypothetical protein; Complete proteome.

SQ SQUINTE 293 AA; 33721 MW; 44971/45949/46964 CPG64

Query March 100.0% Score 38, FR 16, Length 293

Best Local Similarity 66.7% Pred No 2e+02

Matches 4; Conservative 2; Mismatches 3; Indels 0; Gaps 0

QY 1 WXXWHF 6

Db 101 WBPWHF 106

## RESULT 15

Q9EXF1

AC Q9EXF1

DT 01-MAR-2001 (TrEMBLrel 16, Created)

DT 01-MAR-2001 (TrEMBLrel 16, Last sequence update)

DT 01-JUN-2002 (TrEMBLrel 21, Last annotation update)

DE Putative phosphotransferase F stability specific component IIC

DE (Fragment)

GN ATIC

OS Listeria monocytogenes

OC Bacteria; Firmicutes; Bacillus/Clostridium group; Bacillales;

OC Listeriaceae; Listeria

OX NCBI\_TaxID=1629

RN (1)

RP SEQUENCE FROM N.A.

RC STRAIN-1028

RX MEDLINE=2112373; PubMed=11316371

RA Sakiant-onstorgues H., Pontan E., Goossens P.L.,

RT "Characterisation of a Listeria monocytogenes mutant deficient in D-

RL amino acid fermentation."

RL Res. Microbiol. 152:175-177(2001)

DR EMBL: AEO35550; CAC00641.1

DR InterPro: IPR004703; Gal\_spec\_IIC

DR Pfam: PF03611; IIC-GAT\_1

KW Phosphotransferase

FT NON\_TER

FT NON\_TER

SQ SEQUENCE 310 AA; 33721 MW; ACP9P9PC999ACF CPG64

Query March 100.0% Score 39, FR 2; Length 310

Best Local Similarity 66.7% Pred No 2e+02

Matches 4; Conservative 2; Mismatches 3; Indels 0; Gaps 0

QY 1 WXXWHF 6

Db 18 WBYWHF 23

Search completed: February 20, 2003, 10:43:22

Job time: 25.3333 secs



GenCore version 5.1.3  
Copyright 1993 - 2003 Copyright Ltd

OM protein - protein search, using sw model

Run on: February 20, 2003, 16:16:40, Search time: 29:33:56.07s  
(without alignments)  
27.26 Million coll. updates/sec

Title: US-09-912-414-11

Seitens

### Scoring table: PLOCHME200Y

[illegible]

10-11-68

total number of hits satisfying chosen parameters: 90847

Minimum DB seg	length = 0
Maximum DB seg	length = 1000

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Maximum March 1998

[illegible][illegible]

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
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8	/S1D2/gcgatA/geneSeq/geneScp_emb1/AA1997.DAT
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20	/S1D2/gcgatA/geneSeq/geneScp_emb1/AA1999.DAT
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	38	100.0	6	21	AA801492	Peptide which binds
2	38	100.0	6	21	AA801499	Peptide which binds
6	38	100.0	6	21	AA801505	Peptide which binds
4	38	100.0	6	21	AA801506	Peptide which binds
5	38	100.0	6	22	AB840893	Peptide H8199 encodes
6	38	100.0	6	22	AAW61753	Human brain expressed
7	38	100.0	6	22	AAW74549	Human bone marrow
8	38	100.0	4	22	AAW34763	Peptide H8703 encodes
9	38	100.0	4	22	AAW14763	Peptide H8703 encodes
10	38	100.0	4	24	AA024892	Human albumin fusion

[illegible]

```

XX  Claim 6; Page 2, 42pp; English.
XX
XX  Peptides which bind to the DNA binding domain of transcription
XX  factor E2F and inhibit cell cycle progression may be useful as
XX  research agents to investigate the interaction between E2F and DP-1,
XX  or the activation of transcription by E2F/DP-1 heterodimers. They
XX  may also be used for inducing apoptosis and/or cell cycle arrest in
XX  a cell, particularly for treatment of cancer or other proliferative
XX  disorders such as psoriasis and restenosis.
XX
SQ  Sequence 6 AA:
XX
XX  Query Match: 100.0%, Score 38, DB 21, Length 6,
XX  Best Local Similarity: 66.6%, Pos: 10, 7, 100.0%,
XX  Mismatch 4, Conservative 2, Mismatches 0, Indels 0, Gaps 0,
XX
QY  1 WXXWHF 6
XX  |||||
XX  : WXPWHF 6
XX
XX  RESULT 2
XX  ID AAB01499 standard; peptide; 6 AA
XX  AC AAB01499;
XX  DE 08-NOV-2000 (first entry)
XX
XX  Peptide which binds to transcription factor E2F 1 DNA binding domain
XX
XX  DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1,
XX  activation, transcription, apoptosis, proliferation, disorder,
XX  psoriasis, restenosis.
XX
XX  Synthesis:
XX
XX  KQ: Transcription Factor
XX  FT Misc difference 2
XX  FT Misc difference 3
XX  FT Misc difference 3 /note="Any amino acid"
XX
XX  W0200044771-A1.
XX  PD 03-AUG-2000.
XX
XX  de DNA binding domain domain
XX  PR 26-NOV-1999 2000-0001710
XX  PA (PROL-) PROLIFIX LTD.
XX
XX  Mueller E, Kobermann HE, Montigiani S;
XX
XX  WFI, 2000 512806/48
XX
XX  Peptides binding to the DNA binding domain of transcription factor E2F
XX  and inhibiting cell cycle progression, useful for the treatment of
XX  cancer
XX
XX  Claim 4; Page 9, 42pp; English.
XX
XX  Peptides which bind to the DNA binding domain of transcription
XX  factor E2F and inhibit cell cycle progression may be useful as
XX  research agents to investigate the interaction between E2F and DP-1,
XX  or the activation of transcription by E2F/DP-1 heterodimers. They
XX  may also be used for inducing apoptosis and/or cell cycle arrest in
XX  a cell, particularly for treatment of cancer or other proliferative
XX  disorders such as psoriasis and restenosis.
XX
SQ  Sequence 6 AA:

```

```

XX  Query Match: 100.0%, Score 38, DB 21, Length 6,
XX  Best Local Similarity: 66.6%, Pos: 10, 7, 100.0%,
XX  Mismatch 4, Conservative 2, Mismatches 0, Indels 0, Gaps 0,
XX
QY  1 WXXWHF 6
XX  |||||
XX  : WXPWHF 6
XX
XX  RESULT 3
XX  ID AAB01505 standard; peptide; 6 AA.
XX  AC AAB01505;
XX  DE 08-NOV-2000 (first entry)
XX
XX  Peptide which binds to transcription factor E2F 1 DNA binding domain.
XX
XX  DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1,
XX  activation, transcription, apoptosis, proliferation, disorder,
XX  psoriasis, restenosis.
XX
XX  Synthesis:
XX
XX  W0200044771-A1.
XX  PD 03-AUG-2000.
XX
XX  de DNA binding domain domain
XX  PR 26-NOV-1999 2000-0001710
XX  PA (PROL-) PROLIFIX LTD.
XX
XX  Mueller E, Kobermann HE, Montigiani S;
XX
XX  WFI, 2000 512806/48.
XX
XX  Peptides binding to the DNA binding domain of transcription factor E2F
XX  and inhibiting cell cycle progression, useful for the treatment of
XX  cancer
XX
XX  Claim 4; Page 26, 42pp; English.
XX
XX  Peptides which bind to the DNA binding domain of transcription
XX  factor E2F and inhibit cell cycle progression may be useful as
XX  research agents to investigate the interaction between E2F and DP-1,
XX  or the activation of transcription by E2F/DP-1 heterodimers. They
XX  may also be used for inducing apoptosis and/or cell cycle arrest in
XX  a cell, particularly for treatment of cancer or other proliferative
XX  disorders such as psoriasis and restenosis.
XX
SQ  Sequence 6 AA:
XX
XX  Query Match: 100.0%, Score 38, DB 21, Length 6,
XX  Best Local Similarity: 66.6%, Pos: 10, 7, 100.0%,
XX  Mismatch 4, Conservative 2, Mismatches 0, Indels 0, Gaps 0,
XX
QY  1 WXXWHF 6
XX  |||||
XX  : WXPWHF 6
XX
XX  RESULT 4
XX  ID AAB01506
XX  AC AAB01506 standard; peptide; 6 AA.
XX  AC AAB01506;
XX  DE 08-NOV-2000 (first entry)
XX
XX  Peptide which binds to transcription factor E2F 1 DNA binding domain.
XX
XX  DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1,
XX  activation, transcription, apoptosis, proliferation, disorder,
XX  psoriasis, restenosis.
XX
XX  Synthesis:
XX
XX  W0200044771-A1.
XX  PD 03-AUG-2000.
XX
XX  de DNA binding domain domain
XX  PR 26-NOV-1999 2000-0001710
XX  PA (PROL-) PROLIFIX LTD.
XX
XX  Mueller E, Kobermann HE, Montigiani S;
XX
XX  WFI, 2000 512806/48.
XX
XX  Peptides binding to the DNA binding domain of transcription factor E2F
XX  and inhibiting cell cycle progression, useful for the treatment of
XX  cancer
XX
XX  Claim 4; Page 26, 42pp; English.
XX
XX  Peptides which bind to the DNA binding domain of transcription
XX  factor E2F and inhibit cell cycle progression may be useful as
XX  research agents to investigate the interaction between E2F and DP-1,
XX  or the activation of transcription by E2F/DP-1 heterodimers. They
XX  may also be used for inducing apoptosis and/or cell cycle arrest in
XX  a cell, particularly for treatment of cancer or other proliferative
XX  disorders such as psoriasis and restenosis.
XX
SQ  Sequence 6 AA:

```



DE Peptide which binds to transcription factor E2F-1 DNA binding domain  
 XX  
 XX DNA binding, transcription factor E2F-1, E2F-1, p21, cyclin, p1,  
 KW activation, transcription, apoptosis, proliferative disorder,  
 XX psoriasis, restenosis.  
 XX  
 OS Synthetic.  
 PN WO200044771-A1.  
 XX  
 PD 03-AUG-2000.  
 XX  
 PF 26-JAN-2000; 2000WO-GR00227.  
 XX  
 PR 26-JAN-1999; WAGR-0001710.  
 XX  
 PA (PROL-) PROLIFIX LTD.  
 XX  
 PI Mueller R, Kontermann PF, Montijani S;  
 XX WPI; 2000-51206/48  
 DR  
 XX  
 PT Peptides binding to the DNA binding domain of transcription factor E2F  
 PT and inhibiting cell cycle progression, useful for the treatment of  
 PT cancer  
 XX  
 XX Example; Page 26; 42pp, English.  
 XX  
 CC Peptides which bind to the DNA binding domain of transcription  
 CC factor P2F and inhibit cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC or the activation of transcription by E2F-1/DP-1 heterodimers. They  
 CC may also be used for inducing apoptosis and/or cell cycle arrest in  
 CC a cell, particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis  
 CC  
 SQ Sequence 6 AA;  
 XX  
 Query Match 100.0%; Score 18; DB 21; Length 6;  
 Best Local Similarity 66.7%; Pred. No. 7.8e+05;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWHF 6  
 DB 1 WVAWHF 6  
 XX  
 RESULT 5  
 ABB40893  
 ID ABB40893 standard; Peptide; 36 AA.  
 XX  
 AC ABB40893;  
 XX  
 DT 04-FEB-2002 (first entry)  
 XX  
 DE Peptide #8399 encoded by human foetal liver single exon probe  
 XX  
 KW Human; foetal liver; gene expression, single exon nucleic acid probe  
 XX  
 OS Homo sapiens.  
 PN WO200157277-A2.  
 XX  
 PD 09-AUG-2001.  
 XX  
 PF 30-JAN-2001; 2001WO-US00669.  
 XX  
 PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 27-SEP-2000; 2000US-0236359.

PF 04-OCT-2000; 2000GP-0024061  
 XX  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 XX  
 PI Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX  
 DR WPI; 2001 483447/52.  
 XX  
 PT Human genome-derived single-exon nucleic acid probes useful for  
 PT analyzing gene expression in human fetal liver -  
 XX  
 PS Claim 27; SEQ ID NO 33528, 639PP + sequence listing, English.  
 CC  
 CC The invention relates to a single-exon nucleic acid probe for  
 CC measuring human gene expression in a sample derived from human foetal  
 CC liver. The single exon nucleic acid probes may be used for predicting,  
 CC measuring and displaying gene expression in samples derived from human  
 CC foetal liver. The present sequence is a peptide encoded by a single exon  
 CC nucleic acid probe of the invention.  
 CC Note: The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from WIPO at ftp://wipo.int/pub/published\_pat\_sequences  
 CC  
 SQ Sequence 36 AA;  
 XX  
 Query Match 100.0%; Score 38; DB 22; Length 36;  
 Best Local Similarity 66.7%; Pred. No. 56;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWHF 6  
 DB 19 WRAWHF 24  
 XX  
 RESULT 6  
 AAM61753  
 ID AAM61753 standard; Protein; 36 AA.  
 XX  
 AC AAM61753;  
 XX  
 DT 01-MAY-2001 (first entry)  
 XX  
 DE Human brain expressed single-exon probe encoded protein SEQ ID NO: 11854.  
 XX  
 KW Human; brain expressed exon; gene expression analysis; probe;  
 KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;  
 KW epilepsy; cancer.  
 XX  
 OS Homo sapiens.  
 PN WO200157275-A2.  
 XX  
 PD 09-AUG-2001.  
 XX  
 PF 30-JAN-2001; 2001WO-US00667.  
 XX  
 PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GP-0024061.  
 XX  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 PI Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX  
 DR WPI; 2001 483446/52.  
 XX  
 PT Single exon nucleic acid probes for analyzing gene expression in human  
 PT brains -

PS Example 4; SEQ ID NO: 33858, 650bp + Sequence Listing: English.

XX The present invention provides a number of single exon nucleic acid  
CC probes which are derived from genomic sequences expressed in the human  
CC brain. They can be used to measure gene expression in brain cell samples,  
CC which may include the hippocampus and cerebral cortex of nervous system,  
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,  
CC epilepsy, and autism. The present sequence is a protein encoded by one of  
CC the probes of the invention.

XX Sequence 36 AA;

Query Match 100.0%; Score 38; DB 22; Length 36;

Best Local Similarity 65.7%; Prod No. 56;

Matches 4; Conservation 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHP 6  
DB 19 WRAWHF 24

RESULT 7  
AAM74549  
ID AAM/4549 standard; protein; 36 AA.

XX AAM74549

XX AAM74549 (first entry)

XX Homo sapiens

XX Homo sapiens

XX Homo sapiens

XX Homo sapiens

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XX Homo sapiens

XX Homo sapiens

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QY 1 WXXWHP 6  
DB 19 WRAWHF 24

RESULT 8  
AAM74549  
ID AAM/4549 standard; protein; 36 AA.

XX AAM74549

XX AAM74549 (first entry)

XX Homo sapiens

XX Homo sapiens

XX Homo sapiens

XX Homo sapiens

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XX Homo sapiens

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XX Homo sapiens

XX Bone marrow; diagnostic; therapeutic; gene therapy; antigenic;  
 KW haematopoiesis; myeloid; lymph cell disorder; tissue regeneration;  
 KW wound healing; nutritional supplement; immune disorder;  
 KW severe combined immunodeficiency; SCID.  
 OS Homo sapiens.  
 XX  
 PN MO200157187-A2.  
 XX  
 PD 09-APR-2001.  
 XX  
 PF 05-FEB-2001; 2001MO US033782.  
 XX  
 PR 03-FEB-2000; 2000US-0496914.  
 PR 20 JUN 2000; 2000US-0598725.  
 PR 19 JUL-2000; 2000US-0620325.  
 PR 10-NOV-2000; 2000US-0700681.  
 PA (HVSF-) HVSFQ INC.  
 XX  
 PI Ford JE, Boyle P, Tang YL, Liu C, Aschell V, Clark F, Xie AJ,  
 PI Ren F, Dimaane RT;  
 DR WPI: 2001-488475/53  
 DR N Filer; AAC22044  
 XX  
 PT Nucleic acids encoding bone marrow polypeptides, useful in diagnostic  
 PT and gene therapy -  
 PS  
 PS Claim 10; Page 248; 392pp; English.  
 CC AAU14602-AAU14794 represent novel bone marrow polypeptides of the  
 CC invention. The proteins and corresponding coding sequences may be used  
 CC in the prevention, diagnosis and treatment of diseases associated with  
 CC inappropriate bone marrow polypeptide expression. For example, to treat  
 CC disorders associated with decreased expression by rectifying mutations  
 CC or deletions in a patient's genome that affect the activity of the  
 CC polypeptides by expressing inactive proteins or to supplement the  
 CC patient's own production of the polypeptide. Additionally, the nucleic  
 CC acids may be used to produce the polypeptides, by inserting the nucleic  
 CC acids into a host cell and culturing the cell to express the protein.  
 CC The nucleic acid and its complementary sequences may also be used as DNA  
 CC probes in diagnostic assays to detect and quantitate the presence of  
 CC similar nucleic acid sequences in samples, and therefore which patients  
 CC may be in need of restorative therapy. The proteins may also be used as  
 CC antigens in the production of antibodies against bone marrow proteins  
 CC and in assays to identify modulators of their expression and activity.  
 CC The anti-bone marrow protein antibodies and antagonists may also be used  
 CC to down regulate expression and activity. The antibodies may also be used  
 CC as diagnostic agents for detecting the presence of the protein in samples  
 CC (e.g. by enzyme linked immunosorbent assay (ELISA)). The proteins  
 CC may be used to regulate haematopoiesis activity, and consequently in the  
 CC treatment of myeloid or lymph cell disorders; in tissue regeneration,  
 CC such as wound healing; as a nutritional supplement; and in treatment of  
 CC immune disorders such as severe combined immunodeficiency (SCID).  
 XX  
 SQ Sequence 43 AA;  
 Query Match 100.0%; Score 38; DP 22; Length 43;  
 Best Local Similarity 66.7%; Pred. No. 67;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWHF 6  
 DB 14 WLIMHF 19  
 RESULT 10  
 ABG64892  
 ID ARG64892 standard; Protein; 53 AA.  
 XX  
 AC ARG64892;

XX  
 DT 27 AUG-2002 (first entry)  
 XX  
 DE Human albumin fusion protein #1567.  
 XX  
 KW Albumin fusion protein; therapeutic protein X; human albumin; HA;  
 KW human serum albumin; HSA; cancer; reproductive disorder;  
 KW digestive disorder; immune disorder; endocrine disorder;  
 KW haematopoietic disorder; neural disorder; connective disorder;  
 KW cytostatic; antiinfectivity; antiinflammatory; antitumor;  
 KW immunomodulator; anti HIV; antidiabetic; haemostatic; ectopic;  
 KW neuroprotective; antiparkinsonian; antimicrobial; neuroleptic;  
 KW osteoprotectic; antiarthritic.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 PN WO200177137-A1.  
 XX  
 PD 18-OCT-2001.  
 XX  
 PF 12-APR-2001; 2001MO US11368.  
 XX  
 PR 12-APR-2000; 2000US-229358P.  
 PR 25 APR-2000; 2000US-199384P.  
 PR 21 FEB-2000; 2000US-266911P.  
 XX  
 PA (HUMAN) HUMAN GENOME SCI INC.  
 XX  
 PI Rosen CA, Haseltine WA;  
 XX  
 DR WPI: 2002-010886/01.  
 XX  
 PT New fusion protein for treating disease e.g. diabetes comprises an  
 PT albumin fused to a therapeutic protein -  
 PS  
 PS Claim 1; Page 1592; 210pp; English.  
 CC The present invention relates to albumin fusion proteins comprising a  
 CC therapeutic protein X and human albumin (HA), also known as human serum  
 CC albumin (HSA). The proteins are useful for treating a disease or  
 CC disorder that may be modulated by therapeutic protein X. The albumin  
 CC extends the shelf-life of protein X, and may increase its biological  
 CC activity in vivo activity. The protein is useful for treating and  
 CC diagnosing disorders such as cancer, reproductive disorders, live  
 CC disorders (e.g. Crohn's disease, ulcerative colitis), immune disorders  
 CC (e.g. acquired immunodeficiency syndrome, AIDS), endocrine disorders  
 CC (e.g. diabetes), haematopoietic disorders, neural disorder,  
 CC (e.g. Alzheimer's, Parkinson's, Creutzfeldt-Jacob disease,  
 CC encephalomyelitis, meningitis, schizophrenia), and connective disorders  
 CC (e.g. osteoporosis, arthritis). AB053224-AB055518 represent albumin  
 CC fusion proteins of the invention.  
 XX  
 SQ Sequence 53 AA;  
 Query Match 100.0%; Score 38; DP 22; Length 53;  
 Best Local Similarity 66.7%; Pred. No. 81;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWHF 6  
 DB 21 WAGWHF 26  
 RESULT 11  
 AAE21233  
 ID AAE21233 standard; Protein; 53 AA.  
 XX  
 AC AAE21233;  
 XX  
 AC AAE21233;  
 XX  
 DT 01 JUL-2002 (first entry)  
 XX  
 DE Human gene 18 encoded secreted protein HHRM45, SEQ ID NO:98.



XX Sequence 57 AA;  
SQ Query Match 100.0%; Score 38, DB 23, Length 57;  
Best Local Similarity 66.7%; Pred. No. 87;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
OY 1 WXXWHF 6  
|:|:|:|  
DB 4 WPFWHF 9

RESULT 13  
ABP07322  
ID ABP07322 standard; Protein: 65 AA.  
XX  
AC ABP07322;  
XX  
DT 24-JUN-2002 (first entry)  
XX  
DE Human ORFX protein sequence SEQ ID NO:14626.  
XX  
KW Human: Open reading frame; ORFX; gene therapy; cancer; cirrhosis;  
KW hyperproliferative disorder; psoriasis; benign tumour; haemorrhage;  
KW degenerative disorder; osteoarthritis; neurodegenerative disorder;  
KW cardiovascular disease; diabetes mellitus; systemic lupus erythematosus;  
KW hypertension; hypothyroidism; cholesterol ester storage disease;  
KW immune deficiency; immune disorder; infectious disease;  
KW autoimmune disorder; rheumatoid arthritis; autoimmune thyroiditis;  
KW myasthenia gravis.  
XX  
OS Homo sapiens.  
XX  
PN MO200192523-A2.  
XX  
XX 06-DEC-2001.  
XX  
PF 29-MAY-2001; 2001MO-US10836.  
XX  
PP 10-MAY-2000; 2000US-206132P.  
XX 29-MAY-2000; 2000US-206132P.  
XX  
PA (CIBA-) CIBA-CORP.  
XX  
PI Shinkels RA, Leach MD;  
XX  
DR MPI: 2002-106308/14.  
XX N-PSDB: ABN23074.  
XX  
PT Novel human polypeptides and polynucleotides useful for diagnosing,  
PT preventing and treating cardiovascular disease, neurodegenerative,  
PT hyperproliferative disorders and autoimmune disorders -  
XX  
PS Disclosure: SEQ ID 14626; 1037pp; English.

XX The present invention describes substantially purified human proteins  
XX (referred to as open reading frame, ORFX, where X is 1-11491 (see Table 1  
XX in the specification) ABN15762 to ABN27252 encode the human ORFX  
XX protein given in ABP00010 to ABP11500. ORFX proteins are useful for  
XX treating or preventing a pathology associated with an ORFX-associated  
XX disorder in humans, and in the manufacture of a medicament for treating a  
XX syndrome associated with ORFX associated disorder. ORFX polynucleotide  
XX sequences can be used in gene therapy. ORFX sequences can be used in the  
XX treatment of cancer, hyperproliferative disorders, cirrhosis of liver,  
XX psoriasis, benign tumours, keloid, degenerative disorders, haemorrhage,  
XX osteoarthritis, neurodegenerative disorders, disorders related to organ  
XX transplantation, cardiovascular diseases, diabetes mellitus, systemic  
XX lupus erythematosus, hypertension, hypothyroidism, cholesterol ester  
XX storage disease, various immune deficiencies and disorders, infectious  
XX diseases, autoimmune disorders such as multiple sclerosis, rheumatoid  
XX arthritis, autoimmune thyroiditis, myasthenia gravis, graft-versus-host  
XX disease and autoimmune inflammatory eye disease. ORFX proteins are also  
XX useful for treating burns, incisions, ulcers, for treating osteoporosis.

CC bone degenerative disorders, or periodontal disease, and for gut  
CC protection or regeneration and treatment of lung or liver fibrosis,  
CC reperfusion injury in various tissues and conditions resulting from  
CC systemic cytokine damage.  
CC N.R. The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp://ipn.int/pub/published\_pcr\_sequences.  
XX

XX Sequence 65 AA;  
SQ Query Match 100.0%; Score 38, DB 23, Length 65;  
Best Local Similarity 66.7%; Pred. No. 99;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
OY 1 WXXWHF 6  
|:|:|:|  
DB 27 WKSWHF 32

RESULT 14  
AAU56391  
ID AAU56391 standard; Protein: 87 AA.  
XX  
AC AAU56391;  
XX  
DT 27-FEB-2002 (first entry)  
XX  
DE Propionibacterium acnes immunogenic protein #17287.  
XX  
KW SAPHO syndrome, synovitis, acne, pustulosis, hyperostosis, osteomyelitis;  
KW uveitis, endophthalmitis; bone joint; central nervous system; ELISA;  
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
KW dermatological; osteopathic; neuroprotectant.  
XX  
XX  
OS Propionibacterium acnes.  
XX  
PN WO200181581-A2.  
XX  
XX 01-NOV-2001.  
XX  
PF 23-APR-2001; 2001MO-US12865.  
XX  
PP 21-APR-2000; 2000US-199047P.  
XX 02-JUN-2000; 2000US-208841P.  
XX 07-JUL-2000; 2000US-216747P.  
XX  
PA (CORI-) CORIAX CORP.  
XX  
PI Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhattacha A;  
XX L'Alaismeuve J, Zhang Y, Jen S, Carter D;  
XX  
DR MPI: 2001-616774/71.  
XX N-PSDB: AAS59575.  
XX  
PT Propionibacterium acnes polypeptides and nucleic acids useful for  
PT vaccinating against and diagnosing infections, especially useful for  
PT treating acne vulgaris -  
XX  
PS Example 1; SEQ ID No 17586; 1069pp; English.

XX Sequences AAU10105-AAU69017 represent Propionibacterium acnes immunogenic  
XX polypeptides, the proteins and their associated CNA sequences are used in  
XX the treatment, prevention and diagnosis of medical conditions caused by  
XX P. acnes. The disorders include SAPHO syndrome (synovitis, acne,  
XX pustulosis, hyperostosis and osteomyelitis), uveitis and endophthalmitis.  
XX P. acnes is also involved in infections of bone, joints and the central  
XX nervous system, however it is particularly involved in the inflammatory  
XX lesions associated with acne vulgaris. A method for detecting the  
XX presence or absence of P. acnes in a patient comprises contacting a  
XX sample with a binding agent that binds to the proteins of the invention  
XX and determining the amount of bound protein in the sample. The  
XX polypeptides may be used as antigens in the production of antibodies  
XX specific for P. acnes proteins. These antibodies can be used to



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OM Protein - Protein search, using SW method

File: us-09-912-414-11 (without alignments)

48,067 Million cell updates/sec

Title: US-09-912-414-11

Sequence: 1 WXXMHF 6

Scoring table: Gapop 60.0, Gapext 60.0

Searched: 283224 seqs, 96134422 residues

Word size: 0

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post processing: listing first 45 similarities

Database: 1: p1r1  
2: p1r2  
3: p1r3  
4: p1r4

Prod. No. is the number of results predicted by chosen parameters. A score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARY

Result No.	Score	Query Match	Length DB	ID	Description
1	6	100.0	230	270847	hypothetical prote
2	6	100.0	231	271020	hypothetical prote
3	6	100.0	260	271329	hypothetical prote
4	6	100.0	293	283299	hypothetical prote
5	6	100.0	358	1543976	site-specific DNA
6	6	100.0	354	287296	modification methy
7	6	100.0	384	287456	adenine-specific m
8	6	100.0	384	287674	adenine DNA methyl
9	6	100.0	394	287674	putative SAR DNA-b
10	6	100.0	403	287432	site-specific DNA-
11	6	100.0	423	287407	PTS system galacti
12	6	100.0	423	287407	PTS system galacti
13	6	100.0	424	287407	omega 6 desaturase
14	6	100.0	424	287407	omega 6 desaturase
15	6	100.0	439	287407	hypothetical prote
16	6	100.0	444	287407	hypothetical prote
17	6	100.0	447	287407	hypothetical prote
18	6	100.0	448	287407	hypothetical prote
19	6	100.0	452	287407	hypothetical prote
20	6	100.0	452	287407	hypothetical prote
21	6	100.0	462	287407	hypothetical prote
22	6	100.0	472	287407	hypothetical prote
23	6	100.0	475	287407	hypothetical prote
24	6	100.0	487	287407	hypothetical prote
25	6	100.0	492	287407	hypothetical prote
26	6	100.0	497	287407	hypothetical prote
27	6	100.0	497	287407	hypothetical prote
28	6	100.0	504	287407	hypothetical prote
29	6	100.0	508	287407	hypothetical prote

30	6	100.0	508	287407	nucleolar protein
31	6	100.0	511	287407	nucleolar protein
32	6	100.0	511	287407	nucleolar protein
33	6	100.0	550	287407	SAR DNA-binding pr
34	6	100.0	560	287407	SAR DNA-binding pr
35	6	100.0	640	287407	hypothetical prote
36	6	100.0	718	287407	hypothetical prote
37	6	100.0	771	287407	hypothetical prote
38	6	100.0	771	287407	hypothetical prote
39	6	100.0	771	287407	hypothetical prote
40	6	100.0	771	287407	hypothetical prote
41	6	100.0	771	287407	hypothetical prote
42	6	100.0	771	287407	hypothetical prote
43	6	100.0	771	287407	hypothetical prote
44	6	100.0	771	287407	hypothetical prote
45	6	100.0	771	287407	hypothetical prote

## ALIGNMENTS

### RESULT 1

D70847 hypothetical protein P0059 - Mycobacterium tuberculosis (strain H37Rv)

C1Species: Mycobacterium tuberculosis

C1Date: 17-Jul-1998 #sequence\_revision 1 / Jul-1998 #next\_change 20-Jun-2000

C1Accession: D70847

R1C101, S17, Broach, R., Parkhill, J., Garnier, T., Churcher, C., Harris, D., Galloway, S.,

Connor, P., Davies, P., Devlin, R., Feltwell, T., Gentles, S., Hamlin, N., Holroyd, S.,

Rajandream, M.A., Rogers, J., Sanger, K., Skellern, S., Squares, S.,

Nature 393, 537-544, 1998

A1Authors: Sanger, P., Sulston, J., Taylor, K., Whitehead, S., Barrell, B.G.,

Attey, Deciphering the biology of Mycobacterium tuberculosis from its genome

A1Reference number: A10000, M100000000, P100000000

A1Accession: D70847

A1Status: Preliminary, nucleic acid sequence not shown, translation not shown

A1Molecule type: DNA

A1Features: 1..22..220

A1Cross-reference: DB A1000000, DB A1000000, DB A1000000, DB A1000000, DB A1000000

A1Experimental source: strain H37Rv

A1Genetics:

A1Gene: RV0059

A1Function: Mycobacterium tuberculosis hypothetical protein P0059

A1Query Match: 100.0%, Score: 6, CP: 0, Length: 230

A1Best Local Similarity: 66.7%, Prod. No.: 26

A1Matches: 4, Conservative: 2, Mismatches: 0, Indels: 0, Gaps: 0

Q1: 1 WXXMHF 6

Db: 26 WXXMHF 31

### RESULT 2

A10020 hypothetical protein P1054 - Mycobacterium tuberculosis

C1Species: Mycobacterium tuberculosis

C1Date: 14-Jul-1998 #sequence\_revision 14 Aug 1998 #next\_change 20-Jun-2000

C1Accession: A10020

F1Jawaridavasi, V., Swalla, M., Balakrishna, H., Balakrishna, T., Hille, R., Sankaranarayanan, V.,

M., Ohnishi, Y., Tanaka, T., Tanaka, T., Kudo, Y., Yamashita, T., Kusuda, N., Ozaki,

DNA Res. 5, 55-76, 1998

A1Title: Complete sequence and gene organization of the genome of a hyperthermophilic

A1Reference number: A10000, M100000000, P100000000, E100000000

A1Accession: A10020

A1Status: preliminary, nucleic acid sequence not shown, translation not shown

A1Molecule type: DNA

A1Accession: 1..241..241

A1Cross-reference: DB A1000000, DB A1000000, DB A1000000, DB A1000000, DB A1000000

A1Experimental source: strain OT3

A1Note: this accession replaces an interim accession for a sequence replaced by Genbank

A1Genetics:

A1Gene: P1054









GenCore version 5.1.3  
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OM protein - protein search, using sw model

Run on: February 20, 2003, 11:22:17, Search time: 11.333 Seconds  
(without alignments)  
21.958 Million cell updates/sec

Title: US-09-912-414-11

Perfect score: 6  
Sequence: 1 WXXWHF 6

Scoring table: OLGODX  
Gapop 60.0, Gapext 60.0

Searched: 112432 seqs, 4476328 residues

Word size: 0

Total number of hits satisfying chosen parameters: 112432

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database: SwissProt\_40.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No	Score	Query Match	Length	DB ID	Description
1	6	100.0	358	1	MTCL CAUCR
2	6	100.0	376	1	MTCL RHIMR
3	6	100.0	377	1	MTCL RHIMR
4	6	100.0	424	1	MTCL RHIMR
5	6	100.0	424	1	MTCL RHIMR
6	6	100.0	443	1	MTCL RHIMR
7	6	100.0	448	1	MTCL RHIMR
8	6	100.0	479	1	MTCL RHIMR
9	6	100.0	504	1	MTCL RHIMR
10	6	100.0	511	1	MTCL RHIMR
11	6	100.0	529	1	MTCL RHIMR
12	6	100.0	544	1	MTCL RHIMR
13	6	100.0	640	1	MTCL RHIMR
14	6	100.0	718	1	MTCL RHIMR
15	6	100.0	82	1	MTCL RHIMR
16	6	100.0	85	1	MTCL RHIMR
17	6	100.0	95	1	MTCL RHIMR
18	6	100.0	97	1	MTCL RHIMR
19	6	100.0	101	1	MTCL RHIMR
20	6	100.0	103	1	MTCL RHIMR
21	6	100.0	104	1	MTCL RHIMR
22	6	100.0	104	1	MTCL RHIMR
23	6	100.0	104	1	MTCL RHIMR
24	6	100.0	108	1	MTCL RHIMR
25	6	100.0	111	1	MTCL RHIMR
26	6	100.0	111	1	MTCL RHIMR
27	6	100.0	111	1	MTCL RHIMR
28	6	100.0	112	1	MTCL RHIMR
29	6	100.0	112	1	MTCL RHIMR
30	6	100.0	112	1	MTCL RHIMR
31	6	100.0	112	1	MTCL RHIMR
32	6	100.0	112	1	MTCL RHIMR
33	6	100.0	112	1	MTCL RHIMR

## ALIGNMENTS

RESULT 1	ID	MTCL CAUCR	STANDARD:	PRT:	358 AA.
AC	045971				
DT	30-MAY-2000 (Rel. 39, Created)				
DT	16-OCT-2001 (Rel. 40, Last sequence update)				
DE	Modification methylase Cermi (EC 2.1.1.72) (Adenine-specific				
DE	methyltransferase Cermi) (M.Cermi).				
GN	CCPMW OR CCPW OR CCM378.				
OS	Caulobacter crescentus.				
OC	Proteobacteria; alpha subdivision; Caulobacter group;				
OC	Caulobacter.				
OX	NCBI TaxID=155892;				
RN	[1]				
RP	SEQUENCE FROM N A				
RC	STRAIN=CB1N / NA1000;				
FX	MDLINE=94118303; PubMed=829276;				
PA	Zweigler G., Marczynski G., Shapiro L.;				
RT	"A Caulobacter DNA methyltransferase that functions only in the				
RT	predivisional cell."				
RL	J. Mol. Biol. 235:472-485(1994).				
RN	[2]				
RP	SEQUENCE FROM N A				
PC	STRAIN=ATCC 19099 / C15;				
FX	MEDLINE 1173698; PubMed=11259647;				
PA	Norman M.C., Fekshlyum T.V., Laib M.C., Fekshlyum T.V.,				
PA	Elser J., Helgerson J.F., Alvey M.P., Alvey M.P.,				
PA	Patocka J., Nelson M.C., Newton A., Stephens C., Phadke N.D.,				
PA	Deboy R.T., Dodson R.J., Durkin A.S., Gwin M.L., Hall D.H.,				
PA	Kolchay J.F., Smit J., Craven M.B., Phadke N.D., Phadke N.D.,				
PA	Phadke N.D., Tran P., Wolf A., Yamamoto J., Emelanova M.,				
PA	Shapiro L., Shapiro L., Fraser G.M.;				
PT	"Complete genome sequence of Caulobacter crescentus."				
PT	Proc. Natl. Acad. Sci. U.S.A. 98:4116-4120(2001)				
CC	1. FURTHER THIS METHYLASE REGULATES THE TROUBLE STAINING SEQUENCE				
CC	GANCC AND CAUSES SPECIFIC METHYLATION ON A 2 OR BOTH STRANDS.				
CC	CCPM-METHYLASE METHYLATION HAS IMPROVED CELLULAR FUNCTIONS.				
CC	APPEARS TO CONTRIBUTE TO THE ALTERNATE CELL WALL CONTROL OF DNA				
CC	REPLICATION AND CELLULAR MORPHOLOGY.				
CC	1- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + DNA adenine = S-				
CC	adenosyl-L-homocysteine + DNA 6-methylaminopurine.				
CC	This SwissProt entry is "incomplete" because a substitution				
CC	between the Swiss Institute of Bioinformatics and the EMBL outstation				
CC	the European Bioinformatics Institute. There are no restrictions on its				
CC	use by non-profit institutions as long as the copyright is in no way				
CC	modified and this statement is not removed. Usage by and for commercial				
CC	entities requires a license agreement (See <a href="http://www.ebi.ac.uk/infocentre/">http://www.ebi.ac.uk/infocentre/</a>				
CC	or send an email to <a href="mailto:license@ebi.ac.uk">license@ebi.ac.uk</a> ).				
CC	EMBL, CO1032, AAA18913.1, -				
CC	EMBL, AEC05711, AAK22365.1, -				
CC	HSSP: P11409; 1800				
CC	FRPASF, 2233, M.CERM.				
CC	TIGR: CC0378; -				



DR PRINTS: PP00506; D21N6MTFRASE.  
DR PRINTS: PP00508; S21N4MTFRASE.  
DR PRINTS: PP00047; N6-MTFRASE; 1  
KW Transferase, Methyltransferase, DNA replication.  
SQ SEQUNCE 377 AA; 42203 MW; 67709AA;55803SD;CP064

Query Match	100.0%;	Score 6;	DB 1;	Length 377;
Best Local Similarity	66.7%;	Freq. No. 42;		
Matches	4;	Conservative	2;	Mismatches 0;
				Indels 0;
				Gaps 0;

QY	1	WXXWHF	6
		::	
Db	347	WTFWHF	352

RESULT 4	
FD6C SOYBN	
ID	FD6C SOYBN
AC	048658
	STANDARD;
	PRT; 424 AA

DT	01-FEB-1996	(Rel. 3), Created)
DT	01-FEB-1996	(Rel. 33, last sequence update)
DT	15-JUN-2002	(Rel. 41, last annotation update)

glycine max (Soybean) 1499

OC Elavayora: Viridiplantae: Streptophyta: Embryophyta: Tracheophyta:  
 OC Spermatophyta: Magnoliopsida: euembryophytes, rosales  
 OC Apocynaceae: Euphorbia: Euphorbia: Euphorbia  
 OC NCBI\_TaxID=3847;

KN	1
RP	SEQUENCE FROM N A
RC	TISSUE=Seed;

RX MEDLINE=94345008; PubMed 8066133,  
RA Hitz W D, Carlson T J, Booth J  
BA Vaden N G

RT "Cloning of a higher-plant plastid omega-6 fatty acid desaturase cDNA and its expression in a cyanobacterium";

CC -1- PRODUCTION: CHLOROPLAST OMEGA-6 FATTY ACID DESATURASE: INTRODUCES  
CC THE SECOND DOUBLEBOND IN THE BIOSYNTHESIS OF 16:3 AND 18:3 FATTY  
CC ACIDS: INDICATES CONCENTRATIONS

ADOLEN: IMPROVED EFFICIENCY OF PLANT MEMBRANES: IT IS THOUGHT TO USE PERFLUORIN AS AN ELECTRON DOPANT AND TO AGGREGATE ACIDS ESTERIFIED TO CATIONIC SURFACES AND PHOSPHATIDYLGLYCEROL

CC -1- PATHWAY: POLYUNSATURATED FATTY ACID BIOSYNTHESIS:  
CC -1- SUBCELLULAR LOCATION: CHLOROPLAST, MEMBRANE-BOUND (APPROPRIABLE)  
CC -1- DOMAIN: THE HISTIDINE BOX REMAINS MAY CONTAIN THE ACTIVE SITE

AND/OR BE INVOLVED IN METAL ION BINDING.  
SIMILARITY. BELONGS TO THE FATTY ACID DESATURASE FAMILY.

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CC or send an email to [license@ish-sip.ch](mailto:license@ish-sip.ch)  
CC  
CC  
DE EMBL: 129215, AAA50158.1 -

DR InterPro: IP0001225, PA\_desaturase.  
DR Pfam: PFC0487, PA\_desaturase. 1.  
DR ProDom: PD001081, PA\_desaturase. 1

KM	Oxidoreductase; Fatty acid biosyn. basis;
KW	Transit peptide
ET	Chloroplast (low stringency)

FT	TRANS-1	63	CHOLESTEROL 18:1 STEARICITY.
FT	CHAIN	64	OMEGA-6 FATTY ACID DESATURASE.
FT	DOMAIN	165	HISTIDINE BOX-1.
FT	DOMAIN	201	HISTIDINE BOX-2.

FT	DOMAIN	SEQUENCE
1	361	424 AA, 43641 MW, 4F4P5214B1A2000 030644
2	365	
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Query Match      100.0%, Score 6; DB 1; Length 424;
Best Local Similarity 66.7%; Pred. NO. 46;
Matches 4; Conserved 2; Mismatches 0; Gaps 0

```

QY	1	WXXWHF	6
		:	
Db	251	WLMWHF	2

RESULT 5	
Y412_ARATH	
ID Y412_ARATH	STANDARD;
CC 00450	PRT; 439 AA

DT	DT	DT
01-NOV-1997	(P01	35, Created)
01-NOV-1997	(P01	35, last sequence update)
15-JUN-2002	(P01	41, last annotation update)

DE Hypothetical Protein ARSg2120  
GN ATE527120\_QP TM021B04\_12\_QP T21B4\_30  
OS Arabidopsis thaliana (Mouse-ear cress)

00 Equisetata; Viridiplantae; Streptophyta; Embryophyta; Charophyta;  
00 Setaria; Poaceae; Magnoliophyta; eudicotyledons; core eudots; Rosidae;  
00 eunotsids II; Brassicales; Brassicaceae; Arabidopsis.

OX	NCBI TaxID-3702;
RN	[1]
EF	SEQUENCE FROM N.A.

PG STRAIN-CV, Columbia;  
PX MEDLINE=2016721; PubMed=1133714,  
FA Tripathi S., Kaneko T., Nakamura Y., Kikuchi H., Kato T., Asamizu E.,

FA Miyajima N, Sasamoto S, Piyarat T, Higashihata T, Kawashima K,  
FA Kikuchi W, Matsumoto M, Matsuno A, Muraiki A, Iwakayama S,  
PA Miyazaki N, Naruo K, Oyama S, Shirota S, Takeuchi G, Wada T,  
PA

RA Watanabe A., Yamada M., Yasuda M., Sato S., de la Bastide M.,  
PA Huang F., Spiegel L., Gao J., O'Shaughnessy A., Preston P.,  
BA Hobermann K., Murray T., Johnston P., Phillips T., Nelson G.

RA Strongking T, Papin K, Spieth U, Sakon M, Armstrong J, Becker M  
RA Belter E, Cordum H, Cordes M, Courtney B, Courtney W, Dante M,  
RA Du H, Edwards J, Eymann S, Haakensen P, Hamar E, Kattelle P,  
RA

PA Leonard S., Meyer R., Mulvaney E., Ozersky P., Riley A., Stromwater C.  
PA Wajsbort-Welsherson C., William A., Yoakum M., Bell M., Dedia N.,  
PA Parnell L., Shah P., Rodriguez M., Hoon See L., Vill D., Baker J.

PA Kitchhoff V, Toth K, King L, Rahret A, Miller P, Mayra M,  
FA Witt-Lonsdale F, Westoby W P, Wilson R F, Murphy G, Hancock J,  
WA Werger M J, Wardell D, Westoby A, Sclafani W, Dahl T

PA Entlarf P. O., Torgny N., Haller N., Foll E., Jahn M. S.,  
PA Langham S. A., McCallagh E., Robben C., Zygmunt P., Zimmermann W.,  
PA Bannemeyer H., Wedler N., Balke K., Wedler F., Bators C.

PA  
Kampferger G., Weiser H., Balke K., Weiser E., Reuels S.,  
van Staveren M., Dirkse W., Mooijman P., Klein Lankhorst F.,  
Woltzenruegg T., Bothe G., Pose M., Hauf J., Bernolser S., Hempel S.,  
Eilertsen M., Vahedi A., Wittmann R., Schellhorn U.

PA Prof. Dr. habil. W. Langbein, St. Vildriedl R., Gießen U., Arolsen W.  
PA Bents O., Lemcke K., Koljascov G., Mayer K., Fodis S., Schraf H.,  
PA Schrollner C., Garcia P., Mewes H.-W., Bayan M., Eysen T.,

RT "Sequence and analysis of chromosomes of the plant *Arabidopsis thaliana*,"  
PL Nature 409:823-826(2000).

CC 11. SIMILARITY. BELONGS TO THE NOPE/NEPS6 FAMILY.  
CC -----  
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```
CC -----
DR EMBL, AF007271; AAB61073.1; -.
DR InterPro; IPR002697, NFE.
```

DR	Pfam; PF01798; Nop; 1.
DR	ProDom; PD004104; Nop; 1.
KW	Hypothetical protein.

50 SENTENCE 439 AA, 4792 MM, EEEEEE4FE2E41EB CFC64;  
Query Match 100.0% Score 6 MD 1 Length 439.

Best Local Similarity 66.7%; Pred. No. 47;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps

QY 1 WXXMIF 6  
| : | | |



THE SECOND POWERED IN THE BIOSYNTHESIS OF 16:1 AND 18:1 FATTY ACIDS, IMMEDIATE CONSTITUENTS OF PLANT MEMBRANES. IT IS THOUGHT TO USE FERRITIN AS AN ELECTRON DONOR AND TO ACT ON FATTY ACIDS ESTERIFIED TO GALACTOLIPIDS, SULFOLIPIDS AND PHOSPHATIDYLGLYCEROL.

1- PATHWAY: POLYUNSATURATED FATTY ACID BIOSYNTHESIS

2- SUBCELLULAR LOCATION: CHLOROPLAST, MEMBRANE-BOUND (PROBABLY)

3- DEVELOPMENTAL STAGE: HIGHEST LEVELS FOUND IN EXPANDING LEAVES

4- DOMAIN: THE HISTIDINE BOX DOMAIN MAY CONTAIN THE ACTIVE SITE AND/OR BE INVOLVED IN METAL ION BINDING.

5- SIMILARITY: BELONGS TO THE FATTY ACID DESATURASE FAMILY.

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EMBL; U09603; AAA92900.1; -

EMBL; AL022198; CAAB198.1; -

EMBL; AL161579; CAB79813.1; -

EMBL; AT046221; AAK39973.1; -

EMBL; AY058078; AAL24186.1; -

EMBL; AY058078; AAL24240.1; -

InterPro: IPR001225; FA desaturase.

DR Pfam: PF00487; FA desaturase; 1.

DR Pfam: PF001041; FA desaturase; 1.

KM Oxidoreductase; fatty acid biosynthesis; Chloroplast; Membrane;

KM Transit peptide.

FT TRANSIT 1 69 CHLOROPLAST (BY SIMILARITY).

FT CHAIN 70 448 OMEGA-6 FATTY ACID DESATURASE.

FT DOMAIN 171 175 HISTIDINE BOX 1.

FT DOMAIN 207 211 HISTIDINE BOX 2.

FT DOMAIN 367 371 HISTIDINE BOX 3.

SV SEQUENCE 448 AA, 61220 MW, 13672.182kDa; pI=5.74.

Query Match 100.0%; Score 6; DB 1; Length 448;

Best Local Similarity 66.7%; Pred No. 48;

Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHP 6

DB 257 WXXWHP 202

RESULT 8

CATA\_PSEPT STANDARD: PPT; 479 AA

AC 059714;

DT 10-MAY-2000 (rel. 39, Created)

DT 30-MAY-2000 (rel. 39, Last sequence update)

DT 30-MAY-2000 (rel. 39, Last annotation update)

DE Catalase (EC 1.11.1.6).

GN KATA OR CATA.

OS Pseudomonas putida.

OC Bacteria; Proteobacteria; Gamma subdivision; Pseudomonadaceae;

OC Pseudomonas

OC NCBI\_TaxID=303;

PN (1)

RP SEQUENCE FROM N.A.

RC STRAIN=Corvallis;

RA MEDLINE=98019091; PubMed=9358059;

RA Kiv Y.C., Miller C.D., Anderson A.J.;

RT Identification of adjacent genes encoding the major catalase and

RT bacterioferritin from the plant-beneficial bacterium Pseudomonas

RT putida.

RL Gene 199219-224(1997).

CC FUNCTION: FERMENTED HYPEROXYGENATED WATER AIR OXIDATION, SERVES

CC TO PROTECT CELLS FROM THE TOXIC EFFECTS OF HYPEROXYGENATED

CC CATALYTIC ACTIVITY: 3 H<sub>2</sub>O<sub>2</sub> + 2 H<sub>2</sub>O = 2 H<sub>2</sub>O + O<sub>2</sub> (H<sub>2</sub>O<sub>2</sub> + 2 H<sub>2</sub>O = 2 H<sub>2</sub>O + O<sub>2</sub>)

1- SIMILARITY: BELONGS TO THE CATALASE FAMILY.

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EMBL; U63511; AAB8219.1; -

DR HSRP; P42321; CCAE.

DR InterPro: IPR002226; Catalase.

DR Pfam: PF00199; Catalase; 1.

DR PRINTS; PR00067; CATALASE.

DR PROSITE; PS00437; CATALASE\_1; 1.

DR PROSITE; PS00438; CATALASE\_2; 1.

KM Oxidoreductase; Peroxidase; Iron, Heme, Hydrogen Peroxide.

FT ACT SITE 53 53

FT ACT SITE 106 106

FT BINDING 106 106 PROXIMAL HEME LIGAND (BY SIMILARITY).

SV SEQUENCE 479 AA, 53361 MW, 11363.067kDa; pI=5.71

Query Match 100.0%; Score 6; DB 1; Length 479;

Best Local Similarity 66.7%; Pred No. 50;

Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHP 6

DB 209 WXXWHP 214

RESULT 9

SIKI YEAST

IC SIKI YEAST

AC 012460;

DT 01-NOV-1997 (rel. 35, Created)

DT 01-NOV-1997 (rel. 35, Last sequence update)

DT 15-SEP-2002 (rel. 41, Last annotation update)

DE SIKI protein (Nucleolar protein NOP56).

GN SIKI OR NOP56 (Nucleolar protein NOP56).

OS Saccharomyces cerevisiae (Baker's yeast).

OC Eukaryota; Fungi; Ascomycota; Saccharomycetes; Saccharomycetidae; Saccharomycetes.

OC NCBI\_TaxID=4932;

PN (1)

RP SEQUENCE FROM N.A.

RC STRAIN S288C / YPH1;

RA MEDLINE=96040178; PubMed=7547500;

RA Morin P.J., Downs J.A., Snodgrass A.W., Gilmore T.D.;

RT "Genetic analysis of growth inhibition by GATA 1, kappa P alpha in

RT Saccharomyces cerevisiae."

RT Cell Growth Differ. 6:789-798(1995).

RL (2)

RP SEQUENCE FROM N.A.

RC STRAIN=S288C / AB972.

RA Johnston M., Andrews S., Brinkman R., Cooper J., Ding H., Du Z.,

RA Fawell A., Fulton L., Gattung S., Green T., Kieren J., Kucaka T.,

RA Halloworth K., Hawkins D., Hillier L., Jier M., Johnson D.,

RA Johnston L., Langston Y., Latreille P., Mardis E., Meneses S.,

RA Miller N., Nhan M., Pauley A., Peluso D., Rifkin L., Riles L.,

RA Tait S., Trevas E., Vaudin M., Vignati D., Wilcox L., Wilson P.,

RA Wohldman P., Waterson R.;

RT Submitted (SEP-1994) to the EMBL/GenBank/DBPPI databases.

RN (3)

RP CHARACTERIZATION, AND METAGENESIS.

RC MEDLINE=98038077; PubMed=9372940;

RA Gaudier T., Peltier C., Tollervey D., Hurt P.;

RT "Characterization of the yeast protein Pif1 and its role with TopBP1

RT Mol. Cell Biol. 17:7098-7098(1997).

CC FUNCTION: REQUIRED FOR DNA REPLICATION AND REPAIR.









GenCode version 5.1.3  
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OM protein - protein search, using sw model

Run on: February 20, 2003, 11:21:27, Search time: 46 seconds

(without alignments)  
26,876 Million cell writes/sec

Title: US-09-912-414-11

Perfect score: 6

Sequence: 1 WXXWHP 6

Scoring table: OLIGO00X

RefSeq: 20.0, GapPen: 10.0

Searched: 671540 seqs, 20604715 residues

Word size: 0

Total number of hits satisfying chosen parameters: 671540

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summaries

Database:

SPREMBL\_21:  
1: sp\_archaea:  
2: sp\_bacteria:  
3: sp\_fungi:  
4: sp\_human:  
5: sp\_invertebrate:  
6: sp\_mammal:  
7: sp\_mhc:  
8: sp\_organelle:  
9: sp\_phage:  
10: sp\_plant:  
11: sp\_protein:  
12: sp\_virus:  
13: sp\_vertebrate:  
14: sp\_unclassified:  
15: sp\_virus:  
16: sp\_bacteriophage:  
17: sp\_archaea:

Prod No is the number of results provided by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	6	100.0	187	Q9CL54	Q9CL54 pasteurella
2	6	100.0	190	Q9AN05	Q9AN05 unknown homo sapien
3	6	100.0	201	Q9C600	Q9C600 arabidopsis
4	6	100.0	207	Q9AXY2	Q9AXY2 bractea ca
5	6	100.0	213	Q9AXY5	Q9AXY5 brassica na
6	6	100.0	216	Q9AXY7	Q9AXY7 brassica na
7	6	100.0	219	Q9AK47	Q9AK47 arabidopsis
8	6	100.0	222	Q9AXY4	Q9AXY4 arabidopsis
9	6	100.0	240	Q9AL04	Q9AL04 mycobacteri
10	6	100.0	241	Q9AL23	Q9AL23 mycobacteri
11	6	100.0	256	Q9ALN2	Q9ALN2 pyrococcus
12	6	100.0	260	Q9AL25	Q9AL25 arabidopsis
13	6	100.0	264	Q9ALP3	Q9ALP3 triponema p
14	6	100.0	294	Q9AL05	Q9AL05 methanosaic
15	6	100.0	310	Q9EXF1	Q9EXF1 pseudomonas
16	6	100.0	314	Q9AL20	Q9AL20 streptomyces

## ALIGNMENTS

17	6	100.0	381	Q9AT04	Q9AT04 agrobacteri
18	6	100.0	384	Q9AN05	Q9AN05 agrobacteri
19	6	100.0	384	Q9AN05	Q9AN05 guillardia
20	6	100.0	403	Q9AP56	Q9AP56 pyrococcus
21	6	100.0	415	Q9AT04	Q9AT04 thelmodend
22	6	100.0	421	Q9C600	Q9C600 arabidopsis
23	6	100.0	423	Q927H4	Q927H4 listeria m
24	6	100.0	423	Q94Y12	Q94Y12 listeria m
25	6	100.0	423	Q94Y12	Q94Y12 pyrococcus
26	6	100.0	423	Q94Y12	Q94Y12 pyrococcus
27	6	100.0	452	Q927H4	Q927H4 listeria m
28	6	100.0	454	Q927H4	Q927H4 listeria m
29	6	100.0	462	Q927H4	Q927H4 listeria m
30	6	100.0	472	Q94Y12	Q94Y12 listeria m
31	6	100.0	473	Q94Y12	Q94Y12 listeria m
32	6	100.0	473	Q94Y12	Q94Y12 listeria m
33	6	100.0	473	Q94Y12	Q94Y12 listeria m
34	6	100.0	473	Q94Y12	Q94Y12 listeria m
35	6	100.0	487	Q94Y12	Q94Y12 listeria m
36	6	100.0	488	Q94Y12	Q94Y12 listeria m
37	6	100.0	492	Q94Y12	Q94Y12 listeria m
38	6	100.0	492	Q94Y12	Q94Y12 listeria m
39	6	100.0	497	Q94Y12	Q94Y12 listeria m
40	6	100.0	499	Q94Y12	Q94Y12 listeria m
41	6	100.0	508	Q94Y12	Q94Y12 listeria m
42	6	100.0	508	Q94Y12	Q94Y12 listeria m
43	6	100.0	510	Q94Y12	Q94Y12 listeria m
44	6	100.0	510	Q94Y12	Q94Y12 listeria m
45	6	100.0	511	Q94Y12	Q94Y12 listeria m

## RESULT 1

Q9CL54 PESTLIPINAPV; PAT; 187 AA.  
AC Q9CL54;  
DT 01-JUN-2001 (TREMBL) 17; Created)  
DT 01-JUN-2001 (TREMBL) 17; Last sequence update)  
DT 01-JUN-2001 (TREMBL) 20; Last annotation update)  
DE Hypothetical protein PM1135.  
GN PM1135.  
OS Pasteurella multocida.  
OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellales;  
OC Pasteurella.  
OX NCBI\_TaxID=747;  
RN (1)  
RP SEQUENCE FROM N.A.  
RC STEVEN EW73;  
FX MEDLINE:1145866, PubMed 11248100;  
FA May B T, Zhang Q, Li L, Pavittan M, Whitam T, Yip V;  
RT "Complete genomic sequence of Pasteurella multocida PM70";  
PL Proc Natl Acad Sci U S A 94:440-445(1997).  
CC 1. SIMILARITY RELATIONS TO THE TRF/ATP FAMILY OF TRANSCRIPTIONAL  
CC REGULATORS.  
DR EMBL: A606154; AAK03219.1; ;  
DR InterPro: IPR01647; HTM\_TetR.  
DR Pfam: PF09440; TetR\_1.  
DR PRINTS: PR00405; HTHTER.  
KW DNA-binding, Hypothetical protein, Transcription regulation;  
KW Complete proteome.  
KW SEQUENCE 187 AA; 2015 MW; 283EDED14115F CEE4;

Copy Match: 100%; Score: 6; DB: 16; Length: 187;

Best Local Similarity: 66.7%; Prod No: 1.66103;

Matches: 4; Conservative: 2; Mismatches: 0; Gaps: 0;

QY 1 WXXWHP 6  
|||||  
Rb R9 WNIWHP 94

```

Cherry Match          100  0%  00000  0  16  4  00000  00
Beat Lost Slightly   99  0%  00000  0  16  4  00000  00
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0

```

Query Match	Score	EH	Length
Best Local Similarity	78.79	Final RV	1,700,000
Best Local Matrices	4	Conservative	2
		Mismatches	0
		Indels	0
		Gaps	0

Db 156 WNWHP 162

[illegible]

F1 NON-LEK 213 213  
STEFANIE 1971, WM, F0EEF6BDEB06C8C64

Matches 4; 0  
QY 1 WXXXWHP 6



BL Nucleotide position 581-1199.  
 RA SEQUENCE FROM N.A.  
 RA STRAIN COL-100 (JCM8000)  
 RA Fleischmann R.D., Attard R., Brown A.A., Dargatzis D., Miller D.,  
 RA Petersen J., Leber S., Johnson R., Jones M.L., Hall D., Hilgert E.,  
 RA Kolonay J.F., Nelson W.C., Dargatzis D.A., Ermolaeva M.D., Salzberg S.L.,  
 RA Delcher A.L., Ureback T., Weidman D., Kouri H., Gill J., Mikula A.,  
 RA Bressan M., et al. (2001) Genome of *Yersinia enterocolitica* strain 4O8  
 RT "Whole genome sequencing of *Yersinia enterocolitica* strain 4O8  
 RT Laboratory strain"  
 RA STRAIN COL-100 (JCM8000) (EMBL/GenBank/CCDS) JCM8000  
 DR EMBL: A00314281; C0A16240.1  
 DR EMBL: A00314281; A0044357.1  
 DR TIGR: M000007  
 DR Uniprot/SwissProt: P00007  
 RW Hypothetical protein, complete proteome  
 SW SEQUENCE 230 AA; 2556 MW; 1A109D34E-6E9D2AE CRC64

Query Match 100.0% Score 6, DB 16, Length 260  
 Best Local Similarity 66.7% Pred. No. 199402  
 Matches 4, Conservative 2, Mismatches 0, Gaps 0

QY 1 WXXWHF 6  
 DB 26 WXXWHF 21

RESULT 10  
 OS9123 PRELIMINARY: PRT; 241 AA.  
 AC OS9123  
 DT 01-AUG-1998 (TREMblrel\_07, Created)  
 DT 01-AUG-1998 (TREMblrel\_07, Last sequence update)  
 DT 01-MAR-2002 (TREMblrel\_07, Last annotation update)  
 DE Hypothetical protein PH1454.  
 GN PH1454.  
 OS *Pyrobaculum horikoshii*  
 OC Archaea: Euryarchaeota, Thermococci, Thermotogales, Thermotogaceae,  
 OC Pyrobaculus  
 CC NCBI TaxID=53953;  
 OX NCBI TaxID=53953;  
 RN (1)  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-073;  
 RA Fleischmann R.D., Attard R., Brown A.A., Dargatzis D., Miller D.,  
 RA Petersen J., Leber S., Johnson R., Jones M.L., Hall D., Hilgert E.,  
 RA Kolonay J.F., Nelson W.C., Dargatzis D.A., Ermolaeva M.D., Salzberg S.L.,  
 RA Delcher A.L., Ureback T., Weidman D., Kouri H., Gill J., Mikula A.,  
 RA Bressan M., et al. (2001) Genome of *Yersinia enterocolitica* strain 4O8  
 RT "Whole genome sequencing of *Yersinia enterocolitica* strain 4O8  
 RT Laboratory strain"  
 RA STRAIN COL-100 (JCM8000) (EMBL/GenBank/CCDS) JCM8000  
 DR EMBL: A00314281; C0A16240.1  
 DR EMBL: A00314281; A0044357.1  
 DR TIGR: M000007  
 DR Uniprot/SwissProt: P00007  
 RW Hypothetical protein, complete proteome  
 SW SEQUENCE 230 AA; 2556 MW; 1A109D34E-6E9D2AE CRC64

Query Match 100.0% Score 6, DB 16, Length 241  
 Best Local Similarity 66.7% Pred. No. 199402  
 Matches 4, Conservative 2, Mismatches 0, Gaps 0

QY 1 WXXWHF 6  
 DB 193 WXXWHF 198

RESULT 11  
 OS9123 PRELIMINARY: PRT; 236 AA.  
 AC OS9123  
 DT 01-MAR-2001 (TREMblrel\_16, Created)  
 DT 01-MAR-2001 (TREMblrel\_16, Last sequence update)

BL Nucleotide position 581-1199.  
 RA SEQUENCE FROM N.A.  
 RA STRAIN COL-100 (JCM8000)  
 RA Fleischmann R.D., Attard R., Brown A.A., Dargatzis D., Miller D.,  
 RA Petersen J., Leber S., Johnson R., Jones M.L., Hall D., Hilgert E.,  
 RA Kolonay J.F., Nelson W.C., Dargatzis D.A., Ermolaeva M.D., Salzberg S.L.,  
 RA Delcher A.L., Ureback T., Weidman D., Kouri H., Gill J., Mikula A.,  
 RA Bressan M., et al. (2001) Genome of *Yersinia enterocolitica* strain 4O8  
 RT "Whole genome sequencing of *Yersinia enterocolitica* strain 4O8  
 RT Laboratory strain"  
 RA STRAIN COL-100 (JCM8000) (EMBL/GenBank/CCDS) JCM8000  
 DR EMBL: A00314281; C0A16240.1  
 DR EMBL: A00314281; A0044357.1  
 DR TIGR: M000007  
 DR Uniprot/SwissProt: P00007  
 RW Hypothetical protein, complete proteome  
 SW SEQUENCE 230 AA; 2556 MW; 1A109D34E-6E9D2AE CRC64

Query Match 100.0% Score 6, DB 16, Length 256  
 Best Local Similarity 66.7% Pred. No. 29402  
 Matches 4, Conservative 2, Mismatches 0, Gaps 0

QY 1 WXXWHF 6  
 DB 72 WXXWHF 77

RESULT 12  
 OS9250 PRELIMINARY: PRT; 256 AA.  
 AC OS9250  
 DT 01-NOV-1998 (TREMblrel\_08, Created)  
 DT 01-NOV-1998 (TREMblrel\_08, Last sequence update)  
 DT 01-MAR-2002 (TREMblrel\_08, Last annotation update)  
 DE D-D-carboxypeptidase, putative.  
 GN TP0221.  
 OS *Treponema pallidum*  
 OC Bacteria, Spirochaetales, Spirochaetaceae, Treponema  
 CC NCBI TaxID=160;  
 OX NCBI TaxID=160;  
 RN (1)  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-NICHOLS;  
 RA Fleischmann R.D., Attard R., Brown A.A., Dargatzis D., Miller D.,  
 RA Petersen J., Leber S., Johnson R., Jones M.L., Hall D., Hilgert E.,  
 RA Kolonay J.F., Nelson W.C., Dargatzis D.A., Ermolaeva M.D., Salzberg S.L.,  
 RA Delcher A.L., Ureback T., Weidman D., Kouri H., Gill J., Mikula A.,  
 RA Bressan M., et al. (2001) Genome of *Yersinia enterocolitica* strain 4O8  
 RT "Whole genome sequencing of *Yersinia enterocolitica* strain 4O8  
 RT Laboratory strain"  
 RA STRAIN COL-100 (JCM8000) (EMBL/GenBank/CCDS) JCM8000  
 DR EMBL: A00314281; C0A16240.1  
 DR EMBL: A00314281; A0044357.1  
 DR TIGR: M000007  
 DR Uniprot/SwissProt: P00007  
 RW Hypothetical protein, complete proteome  
 SW SEQUENCE 230 AA; 2556 MW; 1A109D34E-6E9D2AE CRC64

Query Match 100.0% Score 6, DB 16, Length 260  
 Best Local Similarity 66.7% Pred. No. 219402  
 Matches 4, Conservative 2, Mismatches 0, Gaps 0

QY 1 WXXWHF 6  
 DB 193 WXXWHF 198

Db 220 WPRWHF 220

## RESULT 13

Q91065 PRELIMINARY PRT 294 AA  
 ID Q91065  
 AC Q91065  
 DT 01-JUN-2002 (TrEMBLrel. 16, last sequence update)  
 DT 01-JUN-2002 (TrEMBLrel. 21, last annotation update)  
 DE Hypothetical protein MA1162.  
 GN MA1162.  
 OS Methanosarcina acetivorans.  
 OC Archaea; Euryarchaeota; Methanococci; Methanosarcinales;  
 CC Methanosarcinaceae; Methanosarcina.  
 OX NCBI\_TaxID=2214;  
 RN [1]  
 RP SPORINTE FROM N.A.  
 RC SPRAIN\_02A / Arco 39395 / DSM 2934.  
 RX MEDLINE 21929763; PubMed 11932318;  
 RA Galagan J E., Nussbaum C., Roy A., Patrizzio M G., MacIntyre J P.,  
 RA Fitzhugh W., Galo S., Fregis P., Smirnov S., Atwood D., Brown A.,  
 RA Allen R., Mayhew M., Wang-Tsai H., Schell J., Wilson R.,  
 RA Lincon L., McKean P., McKernan K., Galanis J., Tittell A., Ye W.,  
 RA Zimmer A., Barber P. D., Cann I., Graham D E., Graham D A., Guss A M.,  
 RA Hedderich P., Ingram-Smith C., Kusterer H C., Kizyoki T A.,  
 PA Leigh J A., Li W., Liu T., Muehlethuy R., Poore J M., Smith K.,  
 PA Springer T A., Tsujimura A., White C., White P H., de Waele E C.,  
 RA Ferry J G., Jarrell K F., Ling H., Macario A L B., Paulsen I.,  
 RA Pritchett M., Sowers K P., Swanson P V., Zinder S H., Zinder F.,  
 RA Metcalf W W., Birren B.;  
 RA "The genome of Methanosarcina acetivorans reveals extensive metabolic  
 RT and physiological diversity";  
 RL Genome Res. 12:432-442(2002)  
 DR EMBL: AEO10783; AAM04583.1;  
 KW Hypothetical protein; Complete proteome;  
 Q2 SEQUENCE 294 AA; 3229 MW; E4210E6E5B2E6 CRO64;

Query Match 100.0%; Score 6; DB 17; Length 289;  
 Best local similarity 66.7%; Pred. No. 2,284,02;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHF 6  
 Db 168 WSIWHF 173

RECURT 14  
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 ID Q91065  
 AC Q91065;  
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)  
 DT 01-MAR-2001 (TrEMBLrel. 16, last sequence update)  
 DT 01-JUN-2002 (TrEMBLrel. 21, last annotation update)  
 DE Hypothetical protein PA2778.  
 GN PA2778.  
 OS Pseudomonas aeruginosa.  
 OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;  
 CC Pseudomonas.  
 OX NCBI\_TaxID=287;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 15692 / PA01;  
 RX MEDLINE=2047337; PubMed=10984043;  
 RA Hickey M J., Britman F S L., Hufnagle W O., Kowalik D T., Lagrou M.,  
 RA Hickey M J., Britman F S L., Hufnagle W O., Kowalik D T., Lagrou M.,  
 RA Garber R L., Goltzy T., Tolentino E., Westlock-Wadman S., Yuan Y.,  
 RA Brody L L., Goltzy T., Tolentino E., Westlock-Wadman S., Yuan Y.,  
 RA Smith K A., Spencer D H., Wong K K S., Wu Z., Paulsen I T.,  
 RA Reizer J., Saiter M H., Hancock P R W., Iori S., Olson M V.;  
 RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an  
 RT opportunistic pathogen";  
 RL Nature 406:959-964(2000).

DR EMBL: AEO31705; AAG06166.1;  
 DP InterPro: IPR005074; Peptidase\_C39  
 DR InterPro: IPR001440; TPR.  
 DR Pfam: PF03412; Peptidase\_C39; 1.  
 KW Hypothetical protein; Complete proteome;  
 Q2 SEQUENCE 293 AA; 32016 MW; DE410E6E5B2E6 CRO64;

Query Match 100.0%; Score 6; DB 15; Length 293;  
 Best local similarity 66.7%; Pred. No. 2,284,02;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHF 6  
 Db 101 WPRWHF 106

RESULT 15  
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 CD C9EXF1 PRELIMINARY FFT 310 AA.  
 AC Q9EXF1;  
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)  
 DT 01-MAR-2001 (TrEMBLrel. 16, last sequence update)  
 DT 01-JUN-2002 (TrEMBLrel. 21, last annotation update)  
 DE Putative phosphotransferase Carabitol specific component 110  
 DE (fragment).  
 GN ATIC.  
 OS Listeria monocytogenes.  
 CC Bacteria; Firmicutes; Bacilli; Clostridium group; Bacillales;  
 CC Listeriaceae; Listeria.  
 OX NCBI\_TaxID=1639;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC SPRAIN-LOC9;  
 RX MEDLINE=2112973; PubMed 11316371;  
 RA Saklani-Josforges H., Fontan E., Goossens P L.;  
 RT "Characterisation of a Listeria monocytogenes mutant deficient in D-  
 RT arabitol fermentation";  
 RL Res. Microbiol. 152:175-177(2001).  
 DR EMBL: AJ292552; CAC20641.1;  
 DR InterPro: IPR004793; Gal\_Spec\_1IC.  
 DR Pfam: PF03611; EITC\_GAT\_1.  
 KW Transferase.  
 FT NON\_TER 1  
 FT NON\_TER 310  
 SQ SEQUENCE 310 AA; 33711 MW; A2F9F92C9C8DACE CRO64;

Query Match 100.0%; Score 6; DB 2; Length 310;  
 Best local similarity 66.7%; Pred. No. 2,284,02;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWHF 6  
 Db 18 WNYWHF 23

Search completed: February 20, 2003, 11:28:57  
 Job time: 46 secs





Genome version 5.1.3  
Copyright 1993-2003 Cambridge Univ

OM protein - protein search, using sw model

February 20, 2003, 11:21:07 : Search time 29.333 seconds  
(without alignment)

27.256 Million cell updates/sec

Title: US-09-912-414-11

Perfect score: 6

Sequence: 1 WXXWHF 6

Scoring table: OLIGOEX

Gapop 60 0 Gapov 60 0

Searched: 909470 seqs 1330520 peptides

Word size: 0

Total number of hits satisfying chosen parameters: 909470

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summaries

Database: A:Geneseq\_101002.\*

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22: /SID52/gcgdata/geneseq/emb1/AA2011.DAT.*
23: /SID52/gcgdata/geneseq/emb1/AA2012.DAT.*
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Pred No is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	6	100.0	6 21 AAB01492	Peptide which bind
2	6	100.0	6 21 AAB01499	Peptide which bind
3	6	100.0	6 21 AAB01505	Peptide which bind
4	6	100.0	6 21 AAB01506	Peptide which bind
5	6	100.0	36 22 AAB040893	Peptide #8399 enco
6	6	100.0	36 22 AAM61753	Human brain expro
7	6	100.0	36 22 AAM74549	Human bone marrow
8	6	100.0	36 22 AAM14663	Peptide #8700 enro
9	6	100.0	43 22 AAM14693	Novel bone marrow
10	6	100.0	43 22 AAB54492	Human albumin fusi

11	6	100.0	53 23 AAB21233	Human gene 18 enro
12	6	100.0	57 23 AAB0179	Protein histone
13	6	100.0	65 23 AAB07322	Human HIF-1 enro
14	6	100.0	87 23 AAB06391	Protein bacterium
15	6	100.0	96 22 AAU14777	Novel bone marrow
16	6	100.0	119 23 AAB06170	Human CFY protein
17	6	100.0	173 23 AAG00884	HIF1 modification
18	6	100.0	173 23 AAG00884	p-paren lipid met
19	6	100.0	175 22 ABB03211	Human secretokole
20	6	100.0	175 22 AAB03371	Human polypptide
21	6	100.0	256 21 AAG00463	Arabis thalia
22	6	100.0	256 21 AAG04075	Arabis thalia
23	6	100.0	256 21 AAG04075	Arabis thalia
24	6	100.0	259 21 AAG04075	Arabis thalia
25	6	100.0	306 21 AAG06810	Arabis thalia
26	6	100.0	306 21 AAG07194	Arabis thalia
27	6	100.0	324 21 AAG05312	Arabis thalia
28	6	100.0	325 21 AAG00462	Arabis thalia
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30	6	100.0	325 21 AAG04074	Arabis thalia
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32	6	100.0	338 21 AAG04074	Arabis thalia
33	6	100.0	376 19 AAG06287	Arabis thalia
34	6	100.0	377 19 AAG06288	Arabis thalia
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42	6	100.0	448 21 AAG047192	Arabis thalia
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45	6	100.0	462 22 AAB05611	Human protein enro

#### ALIGNMENTS

RESULT 1  
AAB01492  
ID AAB01492 standard, peptide: 6 AA.  
XX  
AC AAB01492;  
XX  
UT 68-REV 2000 (first entry)  
XX  
DF Peptide which binds to transcription factor ECF 1 DNA binding domain.  
XX  
PW DNA binding, transcription factor, ECF, ECF 1, cell cycle, DP 1;  
XX  
KW activation, transcription, apoptosis, proliferative disorder;  
XX  
KW peptidase, reesterosis.  
XX  
OS Synthetic.  
XX  
PN WO2000044771-A1.  
XX  
PD 03-AUG-2000.  
XX  
PF 26 JAN 2000; 2000WO/0200227  
XX  
PR 26 JAN 1999; 99GB-0001710.  
XX  
PA (PRO-) PROLIFIX LTD.  
XX  
FI Mueller R, Kontermann RE, Montigiani S;  
XX  
PI WPI, 2000 512806/48.  
XX  
PT Peptide binding to the DNA binding domain of transcription factor ECF 1  
PT and inhibiting cell cycle progression, useful for the treatment of  
PT cancer

[illegible]

QY	1 WXXWHF 6	100.0% Score 6; DB 21; Length 6;
QX		Best Local Similarity 66.67; Std. Dev. 7.86+05;
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DB	1 WXXWHF 6	
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DO	1 WXXWHF 6	
DP	1 WXXWHF 6	
DQ	1 WXXWHF 6	
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EJ	1 WXXWHF 6	
EK	1 WXXWHF 6	
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IF	1 WXXWHF 6	
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II	1 WXXWHF 6	
IJ	1 WXXWHF 6	
IK	1 WXXWHF	

DE Peptide which binds to transcription factor E2F-1 RNA binding domain  
 XX  
 XX DNA binding, transcription factor, E2F, E2F 1, cell cycle, DP-1;  
 KW activation; transcription; apoptosis; proliferative disorder;  
 KW psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 XX  
 PN W0200044771-A1.  
 XX  
 PD 01-AUG-2000.  
 XX  
 PF 26 MAY 2000; 2000US-0608408.  
 XX  
 PP 26-JAN-1999; 99GB-0001710.  
 XX  
 XX (PROL-) PROLIFIX LTD.  
 XX  
 PI Mueller R, Kontermann RE, Montigiani S;  
 XX WPL; 2000-612806/48  
 XX  
 PT Peptides binding to the DNA binding domain of transcription factor E2F  
 PT and inhibiting cell cycle progression, useful for the treatment of  
 PT cancer  
 XX  
 PS Example; Page 26; 42pp; English.  
 XX  
 CC Peptides which bind to the RNA binding domain of transcription  
 CC factor E2F and inhibit cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC or the activation of transcription by E2F-1/DP-1 heterodimers. They  
 CC may also be used for inducing apoptosis and/or cell cycle arrest in  
 CC a cell, particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis.  
 CC  
 SQ Sequence 6 AA;  
 XX  
 Query Match 100.0%; Score 6; DB 21; Length 6;  
 Best Local Similarity 66.7%; Pred. No. 7.8e+05;  
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWHF 6  
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 Db 1 WVAWHF 6  
 XX  
 RESULT 5  
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 ID ABB40893 standard; Peptide; 36 AA.  
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 AC ABB40893;  
 XX  
 DT 04-FEB-2002 (first entry)  
 XX  
 DE Peptide #8399 encoded by human foetal liver single exon probe.  
 XX  
 KW Human; foetal liver; gene expression; single exon nuclear acid probe.  
 XX  
 OS Homo sapiens  
 XX  
 PN W0200157277-A2.  
 XX  
 PD 09-AUG-2001.  
 XX  
 PF 30-JAN-2001; 2001WO-0500669.  
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 PP 04-FEB-2000; 2000US-0180312.  
 XX  
 PR 26-MAY-2000; 2000US-0207456.  
 XX  
 PR 30-JUN-2000; 2000US-0608408.  
 XX  
 PR 03-AUG-2000; 2000US-0612166.  
 XX  
 PR 21-SEP-2000; 2000US-0234687.  
 XX  
 PR 27-SEP-2000; 2000US-0234659.  
 XX

PP 04-DEC-2000; 2000GB-0004263.  
 XX  
 XX (MOLE ) MOLECULAR DYNAMICS INC.  
 XX  
 PI Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX WPL; 2001-483447/52.  
 XX  
 XX Human genome derived single exon nuclear acid probes useful for  
 PT analyzing gene expression in human fetal liver  
 PT  
 XX Claim 27, SEQ ID NO 33528, 639bp + sequence listing, English.  
 XX  
 CC The invention relates to a single exon nucleic acid probe for  
 CC measuring human gene expression in a sample derived from human foetal  
 CC liver. The single exon nucleic acid probes may be used for predicting,  
 CC measuring and displaying gene expression in samples derived from human  
 CC foetal liver. The present sequence is a peptide encoded by a single exon  
 CC nucleic acid probe of the invention.  
 CC Note: The sequence data for this patent did not form part of the  
 CC priority specification, but was obtained in electronic form directly  
 CC from WIPO at ftp.wipo.int/pub/published\_pat\_sequences.  
 CC  
 SQ Sequence 36 AA;  
 XX  
 Query Match 100.0%; Score 6; DB 22; Length 36;  
 Best Local Similarity 66.7%; Pred. No. 25;  
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 |||||  
 Db 19 WRAWHF 24  
 XX  
 RESULT 6  
 AAM61753  
 ID AAM61753 standard; Protein; 36 AA.  
 XX  
 AC AAM61753,  
 XX  
 DT 05-NOV-2001 (first entry)  
 XX  
 DE Human brain expressed single exon probe encoded protein SEQ ID NO: 33854.  
 XX  
 KW Human; brain expressed exon; gene expression analysis; probe;  
 KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;  
 KW epilepsy; cancer.  
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 OS Homo sapiens.  
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 PN W0200157275 A2.  
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 PD 09-AUG-2001.  
 XX  
 PF 30-JAN-2001; 2001WO-0500667.  
 XX  
 PP 04-FEB-2000; 2000US-0190312.  
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 PP 26-MAY-2000; 2000US-0207456.  
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 PR 30-JUN-2000; 2000US-0608408.  
 XX  
 PP 03-AUG-2000; 2000US-0612166.  
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 PP 21-SEP-2000; 2000US-0234687.  
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 PP 27-SEP-2000; 2000US-0234659.  
 XX  
 PP 04-OCT-2000; 2000GB-0024263.  
 XX  
 PA (MOLEF-) MOLECULAR DYNAMICS INC.  
 XX  
 PI Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX WPL; 2001 483446/52.  
 XX  
 FT Single exon nucleic acid probes for analyzing gene expression in human  
 PT brains -  
 XX







XX Sequence 57 AA;  
SQ Query Match 100.0%; Score 6; DB 23; Length 57;  
Best Local Similarity 66.7%; Pred. No. 37;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
QY 1 WXXWHF 6  
|:|:|:|  
Db 4 WFPWHF 9

RESULT 13  
ID ABP07322 standard; Protein; 65 AA.  
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AC ABP07322;  
XX  
DT 24 JUN 2002 (first entry)  
XX  
DE Human CFX protein sequence SEQ ID NO:14626.  
XX  
KW Human; open reading frame; OPRF; gene therapy; cancer; cirrhosis;  
KW hyperproliferative disorder; psoriasis; benign tumor; hemorrhage;  
KW degenerative disorder; osteoarthritis; neurodegenerative disorder;  
KW cardiovascular disease; diabetes mellitus; systemic lupus erythematosus;  
KW hypertension; hypothyroidism; cholesterol ester storage disease;  
KW immune deficiency; immune disorder; infectious disease;  
KW autoimmune disorder; rheumatoid arthritis; autoimmune thyroiditis;  
KW myasthenia gravis  
XX  
OS Homo sapiens  
XX  
PN W0200192523.A2.  
XX  
PD 06-DEC-2001.  
XX  
PF 29-MAY-2001; 2601WO-0810836.  
XX  
PR 30-MAY-2000; 2000US-206132P.  
PR 23-AUG-2000; 2000US-228756P  
XX  
FA (COPR) COPRAX CORP  
XX  
PI Shinkens PA, Leach MD;  
XX  
DR MPI; 2002 106106/14  
XX  
PT N-PSDB; ABN23074.  
XX  
PT Novel human polypeptides and polypeptide-like useful for diagnosing,  
PT preventing and treating cardiovascular disease, neurodegenerative,  
PT hyperproliferative disorders and autoimmune disorders  
XX  
PS Disclosure; SEQ ID 14626; 1037pp; English.

XX The present invention describes substantially purified human proteins  
XX (referred to as open reading frame, OPRF, where X is 1-1449; (see Table 1  
XX in the specification). Applicants to ABP07322 encode the human OPRF  
XX proteins given in ABP0010 to ABP1500. OPRF proteins are useful for  
XX treating or preventing a pathology associated with an OPRF-associated  
XX disorder in humans, and in the manufacture of a medicament for treating a  
XX syndrome associated with OPRF-associated disorder. OPRF polypeptide  
XX sequences can be used in gene therapy. OPRF sequences can be used in the  
XX treatment of cancer, hyperproliferative disorders, cirrhosis of liver,  
XX psoriasis, benign tumors, keloid, degenerative disorders, hemorrhage,  
XX osteoarthritis, neurodegenerative disorders, disorders related to organ  
XX transplantation, cardiovascular diseases, diabetes mellitus, systemic  
XX lupus erythematosus, hypertension, hypothyroidism, cholesterol ester  
XX storage disease, various immune deficiencies and disorders, infectious  
XX diseases, autoimmune disorders such as multiple sclerosis, rheumatoid  
XX arthritis, autoimmune thyroiditis, myasthenia gravis, graft-versus-host  
XX disease and autoimmune inflammatory eye disease. OPRF proteins are also  
XX useful for treating burns, incisions, ulcers, for treating osteoporosis,

XX bone degenerative disorders, or periodontal disease, and for gut  
XX protection or regeneration and treatment of lung or liver fibrosis,  
XX reperfusion injury in various tissues and conditions resulting from  
XX systemic cytokine damage.  
XX N.B. The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from a WFO  
XX at ftp://wipo.int/pub/published\_pat\_sequences.

XX  
SQ Sequence 65 AA;  
QY Query Match 100.0%; Score 6; DB 23; Length 65;  
Best Local Similarity 66.7%; Pred. No. 41;  
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
QY 1 WXXWHF 6  
|:|:|:|  
Db 27 WFSWHF 32

RESULT 14  
ID AAU56391 standard; Protein; 87 AA.  
XX  
AC AAU56391,  
XX  
DT 27-FEB-2002 (first entry)  
XX  
DE Propionibacterium acnes immunogenic protein #17287.  
XX  
KW SAPHO syndrome; synovitis; acne; pustulosis; pyoderma; osteomyelitis;  
KW uveitis; endophthalmitis; bone joint; central nervous system; FUSIA;  
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;  
KW dermatological; osteoporosis; neuroprotectant.  
XX  
OS Propionibacterium acnes.  
XX  
PN W0200191541-A2  
XX  
PD 01-NOV-2001.  
XX  
PF 20-APR-2001; 2001WO-0810845.  
XX  
PR 21-APR-2000; 2000US-199047P.  
PR 02-JUN-2000; 2000US 208841P.  
PR 07-JUL-2000; 2000US 216747P.  
XX  
FA (COPR) COPRAX CORP.  
XX  
PI Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;  
XX  
DR L'Esclapart G, Zhang Y, Sun J, Carter D;  
XX  
DR MPI; 2001-616774/71.  
XX  
PT N-PSDB; AAS59575.  
XX  
PT Propionibacterium acnes polypeptides and nucleic acids useful for  
PT vaccinating against and diagnosing infections, especially useful for  
PT treating acne vulgaris -  
XX  
PS Example 1; SEQ ID NO 17586; 1069pp; English.

XX Sequences AAU91905-AAU6017 represent Propionibacterium acnes immunogenic  
XX polypeptides. The proteins and their associated DNA sequences are used in  
XX the treatment, prevention and diagnosis of medical conditions caused by  
XX P. acnes. The disorders include SAPHO syndrome (synovitis, acne,  
XX pustulosis, hyperostosis and osteomyelitis), uveitis and endophthalmitis.  
XX P. acnes is also involved in infections of bone, joints and the central  
XX nervous system, however it is particularly involved in the inflammatory  
XX lesions associated with acne vulgaris. A method for detecting the  
XX presence or absence of P. acnes in a patient comprises contacting a  
XX sample with a binding agent that binds to the proteins of the invent ion  
XX and determining the amount of bound protein in the sample. The  
XX polypeptides may be used as antigens in the production of antibodies  
XX specific for P. acnes proteins. These antibodies can be used to





GenCore version 5.1.3  
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# OM Protein - Protein search, using SW model

For: 09 February 2003, 10:25:48 / Search time: 24.111 seconds  
(without alignments)  
27.256 Million cell updates/sec

Title: us-09-912-414-9

Perfect score: 31

Sequences: 1 MXXXXF 6

Scoring table:

R/SIM62DX

Gapop: 10.0, Gapext: 0.5

Searched: 000470 seqs, 13050620 residues

Total number of hits satisfying chosen parameters: 328470

Minimum DB seq length: 9

Maximum hit seq length: 20000000

Post processing:

Minimum Match: 49

Maximum Match: 100%

Listing first 45 summaries

Database: A\_Geneseq\_101002.\*

1: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1980.DAT.\*  
2: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1981.DAT.\*  
3: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1982.DAT.\*  
4: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1983.DAT.\*  
5: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1984.DAT.\*  
6: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1985.DAT.\*  
7: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1986.DAT.\*  
8: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1987.DAT.\*  
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11: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1990.DAT.\*  
12: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1991.DAT.\*  
13: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1992.DAT.\*  
14: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1993.DAT.\*  
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16: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1995.DAT.\*  
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18: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1997.DAT.\*  
19: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1998.DAT.\*  
20: /SID52/gcgdata/geneseq/geneeqp-emb1/AA1999.DAT.\*  
21: /SID52/gcgdata/geneseq/geneeqp-emb1/AA2000.DAT.\*  
22: /SID52/gcgdata/geneseq/geneeqp-emb1/AA2001.DAT.\*  
23: /SID52/gcgdata/geneseq/geneeqp-emb1/AA2002.DAT.\*

Prod. No. is the number of results printed by change to have a score greater than or equal to the score of the result being printed, and is derived by analysts of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	31	100.0	6	15	AA057386 Peptide for treati
2	31	100.0	6	15	AA057391 Peptide for treati
3	31	100.0	6	21	AA001492 Peptide which bind
4	31	100.0	6	21	AA001497 Peptide which bind
5	31	100.0	6	21	AA001499 Peptide which bind
6	31	100.0	6	21	AA001505 Peptide which bind
7	31	100.0	6	21	AA001506 Peptide which bind
8	31	100.0	6	21	AA001508 Peptide which bind
9	31	100.0	7	22	AA012181 Polyglutamine repe
10	31	100.0	7	22	AA045777 H11 Binding site c

11	31	100.0	8	15	AA057399 Peptide for treati
12	31	100.0	9	21	AA001498 Peptide which bind
13	31	100.0	9	21	AA001499 Peptide which bind
14	31	100.0	10	15	AA057400 Peptide for treati
15	31	100.0	11	14	AA062149 Haemophilus influenzae
16	31	100.0	11	14	AA062149 Haemophilus influenzae
17	31	100.0	12	15	AA057401 Peptide for treati
18	31	100.0	12	15	AA057401 Peptide for treati
19	31	100.0	12	15	AA057401 Peptide for treati
20	31	100.0	13	18	AA001504 Cystic fibrosis transmembrane
21	31	100.0	13	18	AA001504 Cystic fibrosis transmembrane
22	31	100.0	14	22	AA067760 Human B1A D1 test
23	31	100.0	15	22	AA067760 Human B1A D1 test
24	31	100.0	15	23	AA067760 Human B1A D1 test
25	31	100.0	15	23	AA067760 Human B1A D1 test
26	31	100.0	15	23	AA067760 Human B1A D1 test
27	31	100.0	15	23	AA067760 Human B1A D1 test
28	31	100.0	17	21	AA001491 Peptide for treati
29	31	100.0	19	22	AA067760 Human B1A D1 test
30	31	100.0	20	23	AA067760 Human B1A D1 test
31	31	100.0	22	23	AA067760 Human B1A D1 test
32	31	100.0	22	23	AA067760 Human B1A D1 test
33	31	100.0	22	23	AA067760 Human B1A D1 test
34	31	100.0	22	23	AA067760 Human B1A D1 test
35	31	100.0	22	23	AA067760 Human B1A D1 test
36	31	100.0	22	23	AA067760 Human B1A D1 test
37	31	100.0	22	23	AA067760 Human B1A D1 test
38	31	100.0	22	23	AA067760 Human B1A D1 test
39	31	100.0	22	23	AA067760 Human B1A D1 test
40	31	100.0	22	23	AA067760 Human B1A D1 test
41	31	100.0	22	23	AA067760 Human B1A D1 test
42	31	100.0	22	23	AA067760 Human B1A D1 test
43	31	100.0	22	23	AA067760 Human B1A D1 test
44	31	100.0	22	23	AA067760 Human B1A D1 test
45	31	100.0	22	23	AA067760 Human B1A D1 test

## ALIGNMENTS

RESULT 1

AA057386

ID AA057386 standard; peptide; 6 AA.

XX

AC AA057386;

XX

PT 21-MAP-1995 (first entry)

DE Peptide for treating diseases related to anti-DNA antibodies.

XX

FW Carrier, adsorbing agent, "first map", anti-DNA antibody, the anti-DNA

XX

OS Synthetic.

XX

PM JP06192290-A.

XX

PD 12-000-1994.

XX

PF 18-JAN-1993; 93JP-000609R.

XX

PR 30 SEP 1992, 92JP 026421.

XX

PA (KURS) KURAPAY CO LTD.

XX

WPJ, 1994 060519/22.

XX

PT A peptide and an adsorbing agent, first by inventing it on a

PT carrier - useful for treatment of diseases related to anti-DNA

PT antibodies and immune complexes

XX

PS Disclosure, Page 11; 14pp; Japanese.

XX

CC The sequences given in AA057386 433 are peptides which are all covered

CC by the claimed generic formula:  
 CC H-X (A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q, R, S, T, U, V, W, X, Y, Z)  
 CC A = Trp, Phe or a peptide fragment consisting of 2 residues;  
 CC B = Trp, Phe, Asn or Glu;  
 CC X and Y = a bond or Asp, Glu, Arg, Tyr, His or a peptide fragment  
 CC consisting of 2-10 residues, provided that at least one of  
 CC X or Y are present;  
 CC Z = CH or NH<sub>2</sub>; and  
 CC n = 2-5.  
 CC These peptides may be formulated in a carrier in the preparation of an  
 CC absorbing agent which may be used in the treatment of diseases related  
 CC to anti-DNA antibodies and/or immune complex.  
 CC Sequence 6 AA;  
 CC  
 CC Query Match 100.0%; Score 31; DB 15; Length 6;  
 CC Best Local Similarity 100.0%; First Nov 1990-09-05;  
 CC Matches 3; Conservative 3; Mismatches 0; Gaps 0;  
 CC  
 CC QY 1 WXXXXF 6  
 CC |::|  
 CC Db 1 WXXXXF 6  
 CC  
 CC RESULT 2  
 CC AAR57391 standard; Protein; 6 AA;  
 CC AAR57391;  
 CC 21-MAR-1993 (first entry)  
 CC DE 18-JAN-1993; 31SEP-0000000;  
 CC PF 36-SEP-1993; 92CF-0261803;  
 CC PA (KORS) KIPARAV CO LTD  
 CC WP1, 1994 26010702.  
 CC A peptide not an absorbing agent, but by formulation in an  
 CC carrier useful for treatment of diseases related to anti-DNA  
 CC antibodies and immune complexes  
 CC PS Discovered; Page 11; 14pp; Japanese;  
 CC The sequence given is AAR57391, 430 amino acids, which are all covered  
 CC by the claimed generic formula:  
 CC H-X (A-Bin-Y-Z)  
 CC A = Trp, Phe or a peptide fragment consisting of 2 residues;  
 CC B = Trp, Phe, Asn or Glu;  
 CC X and Y = a bond or Asp, Glu, Arg, Tyr, His or a peptide fragment  
 CC consisting of 2-10 residues, provided that at least one of  
 CC X or Y are present;  
 CC Z = CH or NH<sub>2</sub>; and  
 CC n = 2-5.  
 CC These peptides may be formulated in a carrier in the preparation of an  
 CC absorbing agent which may be used in the treatment of diseases related  
 CC to anti-DNA antibodies and/or immune complex.  
 CC Sequence 6 AA;  
 CC Query Match 100.0%; Score 31; DB 15; Length 6;

CC Best Local Similarity 100.0%; First Nov 1990-09-05;  
 CC Matches 3; Conservative 3; Mismatches 0; Gaps 0;  
 CC  
 CC QY 1 WXXXXF 6  
 CC |::|  
 CC Db 1 WXXXXF 6  
 CC  
 CC RESULT 3  
 CC AAR01497 standard; Peptide; 6 AA;  
 CC AAR01497;  
 CC 08-NOV-2000 (first entry)  
 CC DE Peptide which binds to transcription factor E2F 1 DNA binding domain.  
 CC DNA binding, transcription factor, E2F, E2F1, cell cycle, DP-1,  
 CC activation, transcription, replication, proliferative disorder,  
 CC psoriasis, restenosis.  
 CC OS Synthetic.  
 CC WO200044771-A1.  
 CC 03-AUG-2000.  
 CC 26 JAN 2000; 2000WO 0800227.  
 CC 26 JAN 1993; 1993P 0301730.  
 CC (FISH) FOLIFIX LTD,  
 CC Kallier S, Kallierman PE, Montgomeri G,  
 CC WP1, 2000-532806/48.  
 CC Peptide which binds to the DNA binding domain of transcription factor E2F  
 CC and inhibiting cell cycle progression, useful for the treatment of  
 CC cancer  
 CC Claim of Page 3; 4pp; English.  
 CC Peptide which binds to the DNA binding domain of transcription  
 CC factor E2F and inhibits cell cycle progression may be useful as  
 CC research agents to investigate the interaction between E2F and DP-1,  
 CC or the activation of transcription by E2F1/DP1 heterodimers. They  
 CC may also be used for inducing apoptosis and/or cell cycle arrest in  
 CC a cell particularly for treatment of cancer or other proliferative  
 CC disorders such as psoriasis and restenosis.  
 CC Sequence 6 AA;  
 CC Query Match 100.0%; Score 31; DB 31; Length 6;  
 CC Best Local Similarity 100.0%; First Nov 1990-09-05;  
 CC Matches 3; Conservative 3; Mismatches 0; Gaps 0;  
 CC  
 CC QY 1 WXXXXF 6  
 CC |::|  
 CC Db 1 WXXXXF 6  
 CC  
 CC RESULT 4  
 CC AAR01497 standard; peptide; 6 AA.  
 CC AAR01497;  
 CC 08-NOV-2000 (first entry)  
 CC DE Peptide which binds to transcription factor E2F 1 DNA binding domain.  
 CC



PT Cancer  
 XX  
 XX  
 XX Example, Page 26, 42pp, English.  
 CC Peptides which bind to the DNA binding domain of transcription factor E2F and inhibit cell cycle progression may be useful as research agents to investigate the interaction between E2F and DP-1, or the activation of transcription by E2F-1 in tumor cells. They may also be used for inducing apoptosis and/or cell cycle arrest in a cell, particularly for treatment of cancer or other proliferative disorders such as psoriasis and restenosis.  
 CC  
 AA  
 SQ Sequence 6 AA;

Query Match 100.0%, Score 31, DB 21, Length 6;  
 Best Local Similarity 100.0%, Pval 1.7e-05;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXMXF 6  
 DB 1 WXXMXF 6

RESULT 7  
 AAB01506  
 ID AAB01506 standard, peptide; 6 AA.  
 XX  
 AC AAB01506;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DE Peptide which binds to transcription factor E2F-1 DNA binding domain.  
 XX  
 KW DNA binding; transcription factor; E2F, E2F-1, cell cycle; DP-1, cyclin; transcription; apoptosis; psoriasis; proliferative disorder; psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 XX  
 PN W0200044771-A1.  
 XX  
 PD 03-AUG-2000.  
 XX  
 PE 26-JAN-2000; 2000WC-GP00227.  
 XX  
 PR 26-JAN-1999; 99GB-0001710.  
 XX  
 RA (PROD-) PROLIFIX LTD.  
 XX  
 PI Mueller R, Kottmann RE, Montigiani S;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DE Peptides binding to the DNA binding domain of transcription factor E2F and inhibiting cell cycle progression, useful for the treatment of cancer.  
 XX  
 PS Example, Page 26, 42pp, English.  
 XX  
 CC Peptides which bind to the DNA binding domain of transcription factor E2F and inhibit cell cycle progression may be useful as research agents to investigate the interaction between E2F and DP-1, or the activation of transcription by E2F-1 in tumor cells. They may also be used for inducing apoptosis and/or cell cycle arrest in a cell, particularly for treatment of cancer or other proliferative disorders such as psoriasis and restenosis.  
 CC  
 AA  
 SQ Sequence 6 AA;

Query Match 100.0%, Score 31, DB 21, Length 6;  
 Best Local Similarity 100.0%, Pval 1.7e-05;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXMXF 6  
 DB 1 WXXMXF 6

RESULT 8  
 AAB01508  
 ID AAB01508 standard, peptide; 6 AA.  
 XX  
 AC AAB01508;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DE Peptide which binds to transcription factor E2F-1 DNA binding domain.  
 XX  
 KW DNA binding; transcription factor; E2F, E2F-1, cell cycle; DP-1, cyclin; transcription; apoptosis; psoriasis; proliferative disorder; psoriasis; restenosis.  
 XX  
 OS Synthetic.  
 XX  
 PN W0200044771-A1.  
 XX  
 PD 03-AUG-2000.  
 XX  
 PE 26-JAN-2000; 2000WC-GP00227.  
 XX  
 PR 26-JAN-1999; 99GB-0001710.  
 XX  
 RA (PROD-) PROLIFIX LTD.  
 XX  
 PI Mueller R, Kottmann RE, Montigiani S;  
 XX  
 DT 08-NOV-2000 (first entry)  
 XX  
 DE Peptides binding to the DNA binding domain of transcription factor E2F and inhibiting cell cycle progression, useful for the treatment of cancer.  
 XX  
 PS Example, Page 26, 42pp, English.  
 XX  
 CC Peptides which bind to the DNA binding domain of transcription factor E2F and inhibit cell cycle progression may be useful as research agents to investigate the interaction between E2F and DP-1, or the activation of transcription by E2F-1 in tumor cells. They may also be used for inducing apoptosis and/or cell cycle arrest in a cell, particularly for treatment of cancer or other proliferative disorders such as psoriasis and restenosis.  
 CC  
 AA  
 SQ Sequence 6 AA;

Query Match 100.0%, Score 31, DB 21, Length 6;  
 Best Local Similarity 100.0%, Pval 1.7e-05;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXMXF 6  
 DB 1 WXXMXF 6

RESULT 9  
 AAB01281  
 ID AAB01281 standard, peptide; 7 AA.  
 XX  
 AC AAB01281;  
 XX  
 DT 18-DEC-2001 (first entry)  
 XX  
 DE Polyglutamine repeat domain binding peptide, QPBI-M7FD.  
 XX  
 KW Polyglutamine repeat domain; Huntington; atrophy; ataxin; Huntington receptor; pallidum; pallidum; Huntington disease; Huntington's disease; denervation; pallidum; atrophy; DPLA;  
 XX

KW spinobulbar muscular atrophy; spinocerebellar ataxia; motoric.  
 XX Unidentified.  
 OS  
 XX  
 PN WC200168678-A2.  
 XX  
 PD 20-SEP-2001.  
 XX  
 PF 14-MAR-2001; 2601WC-0509222.  
 XX  
 PR 16-MAR-2000; 2600US 190791P.  
 XX  
 PA (UYDU-) UNITV DIRE.  
 XX  
 PI Strittmatter WL, Burke JR, Nagai Y;  
 XX  
 DE WPI: 2001-414245/71  
 XX  
 PT New polypeptides which polypeptide binding to expanded polyglutamine  
 PT repeat domains, useful for treating neurodegenerative diseases, e.g.  
 PT Huntington's disease.  
 XX  
 PS Example 13; Page 14; 59pp; English  
 XX  
 CC The present sequence is a polyglutamine repeat domain binding peptide.  
 CC The peptide is useful for treating a cell that contains and expresses a  
 CC protein having an expanded polyglutamine region (e.g. huntingtin;  
 CC atrophin 1; ataxin 1; 2, 6 or 7; or antigen receptor protein); or  
 CC treating a neurodegenerative disease characterised by the presence of  
 CC expanded polyglutamine repeats, e.g. Huntington's disease, dentatorubral  
 CC pallidum/lysian atrophy (DRPAL), spinobulbar muscular atrophy, and  
 CC spinocerebellar ataxia types 1, 2, 3, 6 and 7  
 XX  
 SQ Sequence 7 AA;  
 XX  
 Query Match 100.0%; Score 31; DE 22; Length 7;  
 Best Local Similarity 50.0%; Pred. No. 7.8e+05;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWXP 6  
 DB 1 WKWMPF 6  
 XX  
 RESULT 10  
 AAM45777  
 ID AAM45777 standard; Peptide; 7 AA.  
 XX  
 AC AAM45777;  
 XX  
 DT 25-DEC-2001 (first entry)  
 XX  
 DE H11 binding site; cross-reacts with binding peptide (WPI #2049.  
 XX  
 KW Antigen-binding; tumour diagnosis; stress protein peptide complex; SPPC;  
 KW immunogenetically cross-reactive; cancer; immunogenic cancer cell;  
 KW cytotoxicity; tumour specific; immunogenic response inhibitor;  
 KW astrocytoma, fibrosarcoma, myosarcoma, liposarcoma, gliodendrogloma;  
 KW oligodendrogloma; medulloblastoma; primitive neural ectodermal tumour;  
 XX  
 OS Homo sapiens  
 OS Synthetic.  
 XX  
 PN CA229222-A1.  
 XX  
 PD 08-JUN-2001.  
 XX  
 PF 08-DEC-1999; 99CA-2290722.  
 XX  
 PR 08-DEC-1999; 99CA-2290722.  
 XX  
 PA (NOVO) NOVOPIARM BIOTECH INC.  
 XX

PI Kaplan HA, Maiti PK, Fast PG, Herman W, Pan MD, Lewis KE;  
 PI Entwistle JM, MacDonald GC;  
 XX  
 DP WPI: 2001 425937/46.  
 XX  
 PT Composition useful for treating and diagnosing cancer, comprises a stress  
 PT protein-peptide complexes associated with tumor, and isolated  
 PT antigen-binding fragments of an antibody that binds specifically to the  
 PT complex.  
 XX  
 PS Example 4, Page 108, 154pp, English.  
 XX  
 CC The present invention describes a composition (1) comprising stress  
 CC protein-peptide complexes (SPPC) associated with tumors that is  
 CC specifically immunogenetically cross-reactive with cell surface associated  
 CC SPPCs specific to target cancer (TC). Also described is an isolated  
 CC antigen binding fragment of an antibody that binds specifically to SPPCs  
 CC or a population of different SPPCs consisting of immunogenic cancer cell  
 CC surface associated SPPC of TC. (1) has cytotoxic activity and can be  
 CC used in vaccine production and as a tumor-specific immunogenic response  
 CC inducer. (1) is useful for treating 71 types of cancers or tumors in a  
 CC subject, such as astrocytoma, fibrosarcoma, myosarcoma, liposarcoma,  
 CC oligodendrogloma, ependyoma, medulloblastoma, and primitive neural  
 CC ectodermal tumour (PNET). (1) is useful as cancer immunogen including  
 CC vaccines. (1) is useful for diagnostic and palliative use, for detecting  
 CC or imaging cancer cells, and to monitor the course of proliferation of  
 CC malignancy in an individual. AAM43707 to AAM47109 represent peptides  
 CC which are used in the exemplification of the present invention.  
 XX  
 SQ Sequence 7 AA;  
 XX  
 Query Match 100.0%; Score 31; DE 22; Length 7;  
 Best Local Similarity 50.0%; Pred. No. 7.8e+05;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 WXXWXP 6  
 DB 1 WPMWNP 6  
 XX  
 RESULT 11  
 AAR57399  
 ID AAR57399 standard; Protein; 8 AA.  
 XX  
 AC AAR57399;  
 XX  
 DT 21-MAR-1995 (first entry)  
 XX  
 DE Peptide for treating diseases related to anti-DNA antibodies.  
 XX  
 KW Carrier; adsorbing agent; treatment; anti-DNA antibody; immune response;  
 XX  
 OS Synthetic.  
 XX  
 PN JP06192290-A.  
 XX  
 PD 12-JUL-1994.  
 XX  
 PF 18-JAN-1993; 93JP-0006098.  
 XX  
 PR 30-SEP-1992; 92JP-0261921.  
 XX  
 PA (KIPPS) KIPAPAY CO LTD.  
 XX  
 DP WPI: 1994 260510/32.  
 XX  
 PT A peptide and an adsorbing agent prep'd. by immobilising it on a  
 PT carrier useful for treatment of diseases related to anti-DNA  
 PT antibodies and immune complexes  
 XX  
 PS Disclosure; Page 12; 14pp; Japanese.  
 XX  
 CC The sequences given in AAR57386-413 are peptides which are all covered



stimulating factor (G-CSF a haematopoietic growth factor and  
 including evaluation of many factors thought to influence, and be  
 influenced by, production of white blood cells, in the development of  
 components that bind to G-CSF, as reagents for detecting G-CSF receptor  
 or related receptor on living cells, fixed cells, in biological fluid, in  
 tissue homogenates or in purified natural biological materials, in situ  
 staining, fluorescence-activated cell sorting (FACS), Western blotting or  
 enzyme-linked immunosorbent assay (ELISA), in receptor purification or  
 in purifying cells expressing G-CSF on the cell surface (or inside  
 permeabilised cells) as a commercial research reagent for various medical  
 and diagnostic uses or to treat a disease that would benefit from the  
 ability to of a compound to mimic the effects of G-CSF in vivo.  
 The compounds bind specifically to G-CSF and allow for studies of  
 biological activities mediated by the receptor and for the treatment of  
 diseases, disorders and conditions that would benefit from activating or  
 inactivating G-CSF. The present sequence is a G-CSF binding peptide or  
 the invention.

Sequence 9 AA:

Query Match 100.0% Score 31; EB 23; Length 9;

Best Local Similarity 50.0% Pred. No. 7 80.0%;

Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 |||||  
 Db 4 WWCWNF 9

RESULT 14

AARS7400  
 ID AARS7400 standard; Protein; 10 AA

AC AARS7400;

DT 21-MAR-1995 (first entry)

DE Peptide for treating diseases related to anti-DNA antibodies

XX Carrier: absorbing agent, treatment, anti-DNA antibody, immunosuppressant

XX Synthetic

XX JP06192290-A.

XX PD 12 JUL 1994.

XX PF 18-JAN-1993; 93JP-0006098.

XX PR 30-SEP-1992; 92JP 0701921.

XX PA (KUPS) KUPAPAV CO LTD.

XX DR WPI; 1994.360510/12

XX A peptide and an adsorbing agent prep'd by immobilising it on a  
 PT carrier - useful for treatment of diseases related to anti-DNA  
 PT antibodies and immune complexes

XX PS Disclosure; Page 12; 14pp; Japanese

XX The sequences given in AARS7400 are peptides which are all covered  
 CC by the claimed generic formula:

CC H-X-(A-B)n-Y-Z  
 CC A = Trp, Phe or a peptide fragment consisting of 2 residues;

CC B = Trp, Phe, Asn or Glu;

CC X and Y = a bond or Asp, Glu, Arg, Lys, His or a peptide fragment  
 CC consisting of 2-10 residues, provided that at least one of  
 CC X or Y are present;

CC Z = OH or NH<sub>2</sub>; and  
 CC n = 2-5.

CC These peptides may be immobilised on a carrier in the preparation of an

CC absorbing agent which may be used in the treatment of diseases related  
 CC to anti-DNA antibodies and/or immune complex.

Sequence 10 AA:

Query Match 100.0% Score 31; EB 15; Length 10;

Best Local Similarity 50.0% Pred. No. 1 00.0%;

Matches 3; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 WXXWXP 6  
 |||||  
 Db 3 WWCWNF 8

RESULT 15

AAW62149  
 ID AAW62149 standard; Peptide; 11 AA.

AC AAW62149;

DT 16-SEP-1998 (first entry)

DE Haemophilus influenzae tyrosine RNA synthetase binding peptide 22.

XX Identification; ligand; biological activity; target-binding;

XX drug screening; library; inhibitory ligand.

XX Synthetic.

XX Haemophilus influenzae.

XX WO9819162-A1.

XX PD 07-MAY-1998.

XX PF 31-OCT-1997; 97WO-0519638.

XX PP 31-OCT-1996; 96US-0740671.

XX PA (NOVA) NOVALON PHARM CORP.

XX Fowlkes TM, Frolingho JN, Hyde-Fowler PT, Kay BK;

XX WPI; 1998-072389/24.

XX Identifying ligands which mediate biological activity of a protein  
 PT by identifying target-binding ligands and screening a library for  
 PT ligands which inhibit target-binding ligand mediated activity

XX Example 5; Page 100; 14pp; English.

XX A method has been developed for identifying a ligand which mediates the  
 CC biological activity of a target protein (n) by inhibiting the binding  
 CC of (n) to a binding partner. The method comprises: (a) screening a first  
 CC combinatorial library comprising first member ligands for binding to the  
 CC target binding ligands (TBLs), to identifying one or more TBLs; (b)

CC screening a second library comprising second member ligands for the  
 CC ability to inhibit the binding of one or more of the TBLs to the target

CC protein, and so obtaining one or more inhibitory ligands; and (c)

CC determining which of the inhibitory ligands can mediate a biological

CC activity of the target protein. The present sequence represents a

CC potential binding peptide for Haemophilus influenzae tyrosine RNA

CC synthetase from an example of the present invention. The method can be  
 CC used for identifying drugs which can modulate the biological activity of

CC a target protein. It can be used to identify the biological activity of

CC a target protein whose biological function is not known and perhaps  
 CC cannot be determined directly. The method can also be used to identify

CC new inhibitory ligands of specific target proteins. The method provides  
 CC high throughput screens which are essentially identical for similar and  
 CC dissimilar targets, bypassing the need to develop distinct assays for  
 CC biochemically diverse targets.

Sequence 11 AA:

Query Match 100.0%; Score 31; DB 19; Length 11;  
 Best Local Similarity 50.0%; Ered No. 2,1e+02;  
 Matches 3; Conservative 3; Mismatches 0; Indels 0;

QY 1 WXXWXP 6  
 11111  
 Db 5 WPDWGF 10

Search completed: February 20, 2003, 10:47:36  
 CPU time: 31.333 secs